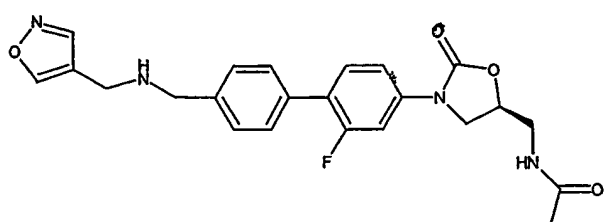
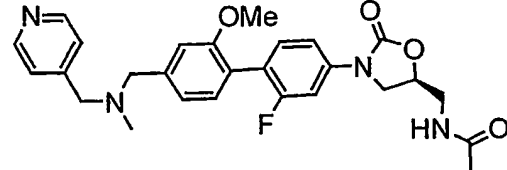
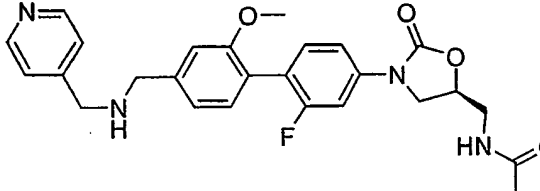
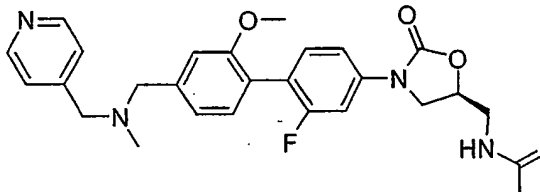
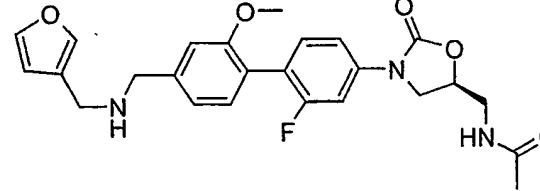
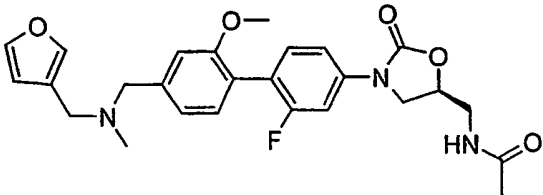
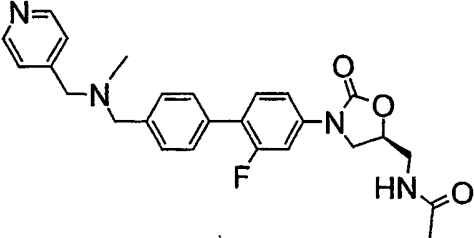
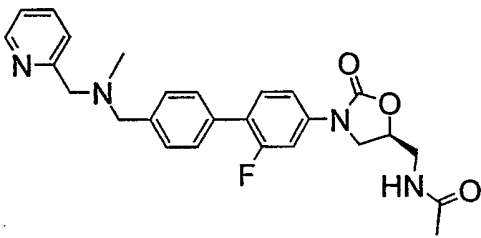
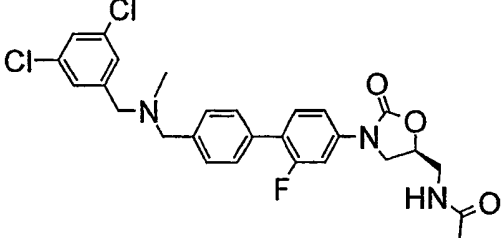
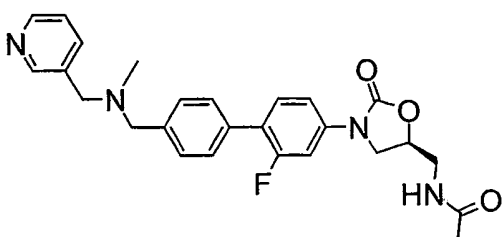
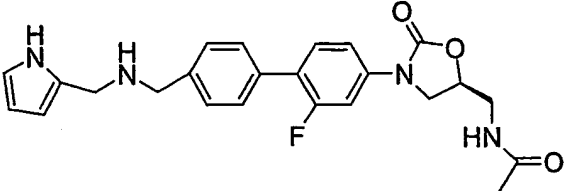
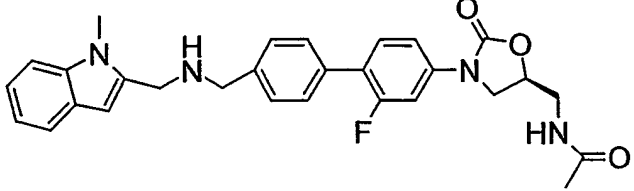
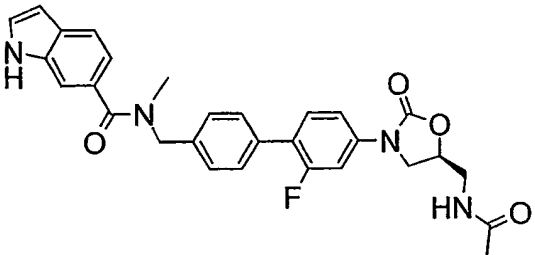
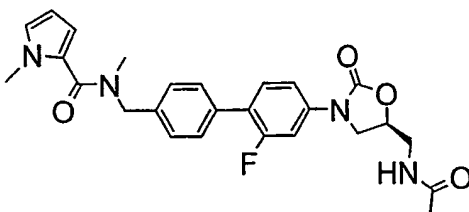
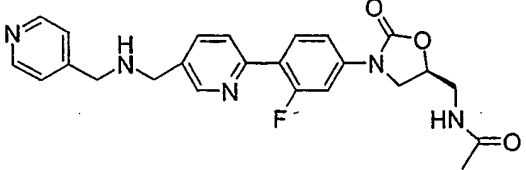
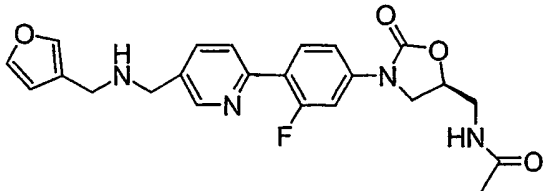
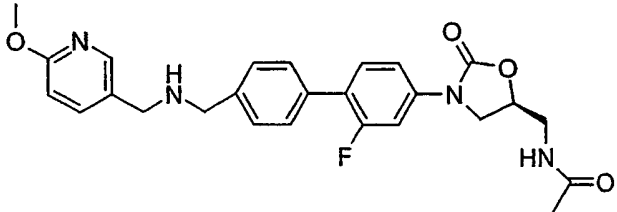
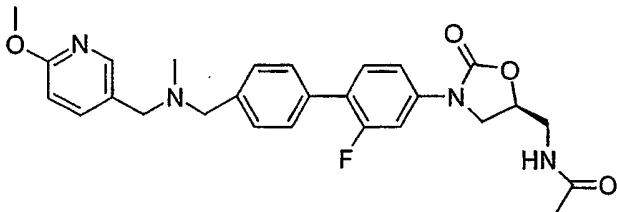
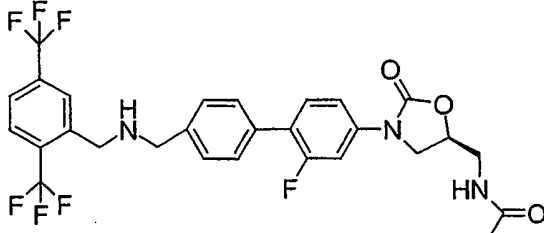
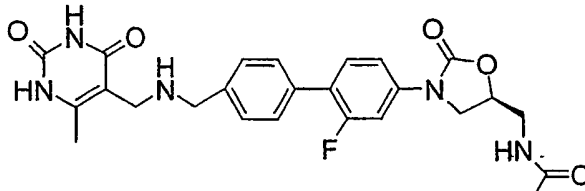
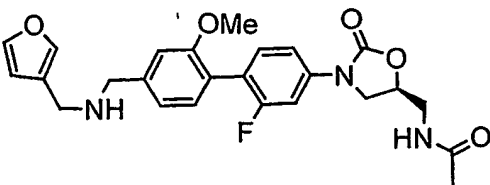
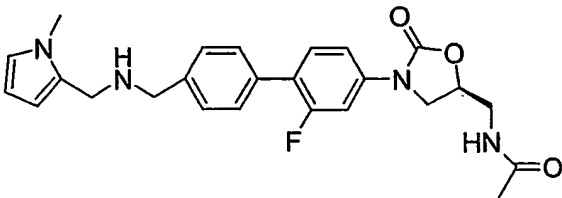
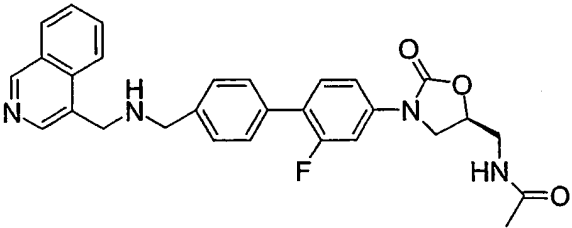
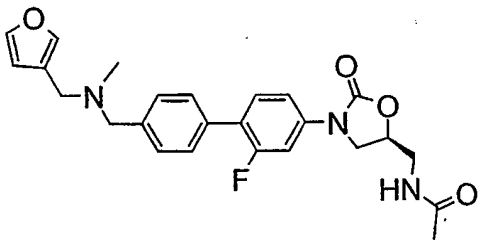
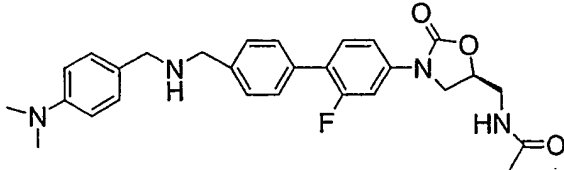


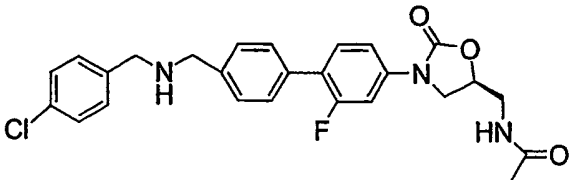
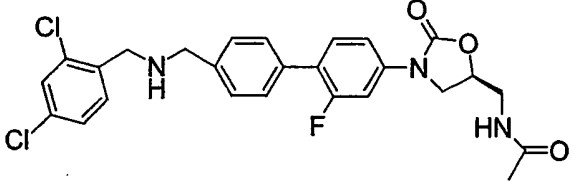
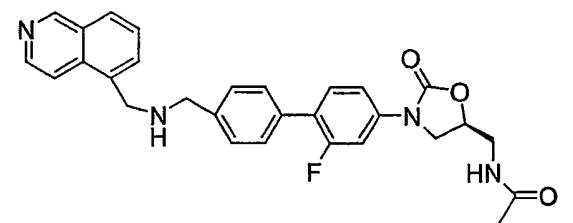
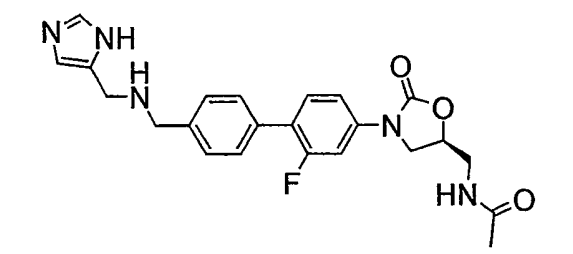
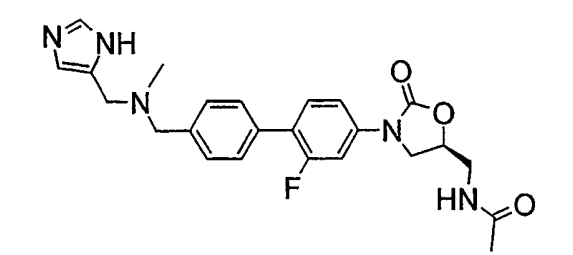
4038	
	N-[3-(2-Fluoro-4'-{[(isoxazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4039	
	N-(3-{2-Fluoro-2'-methoxy-4'-[(methyl-pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4040	
	N-[3-(2-Fluoro-2'-methoxy-4'-{[(pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4041	
	N-(3-{2-Fluoro-2'-methoxy-4'-[(methyl-pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4042	
	N-[3-(2-Fluoro-4'-{[(furan-3-ylmethyl)-amino]-methyl}-2'-methoxy-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

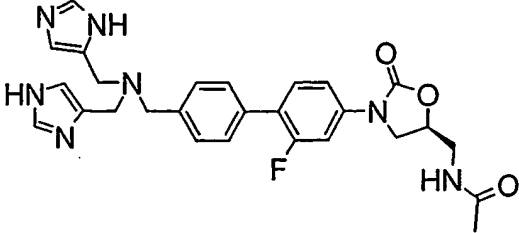
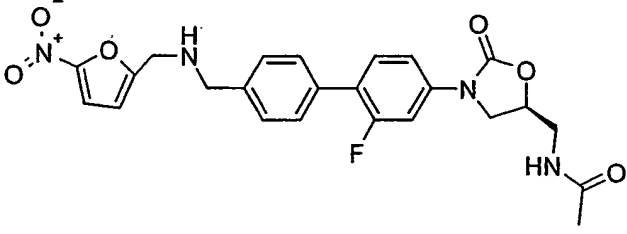
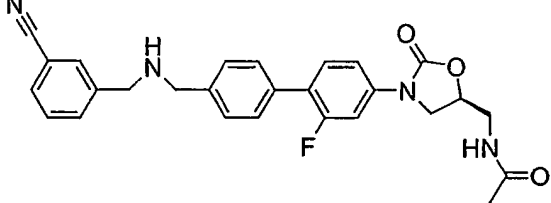
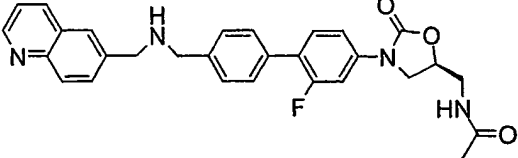
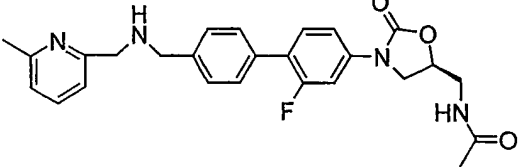
4043	
	N-(3-{2-Fluoro-4'-[(furan-3-ylmethyl-methyl-amino)-methyl]-2'-methoxy-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4044	
	N-(3-{2-Fluoro-4'-[(methyl-pyridin-4-ylmethyl-amino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4045	
	N-(3-{2-Fluoro-4'-[(methyl-pyridin-2-ylmethyl-amino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4046	
	N-[3-(4'-{[(3,5-Dichloro-benzyl)-methyl-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4047	

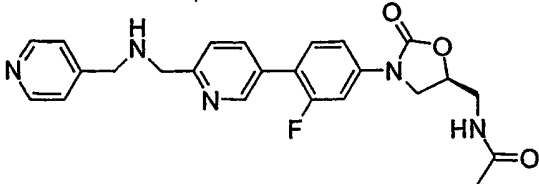
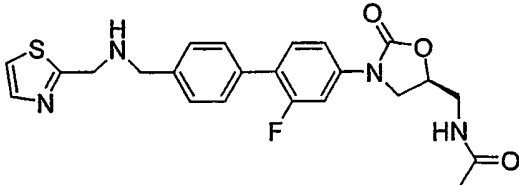
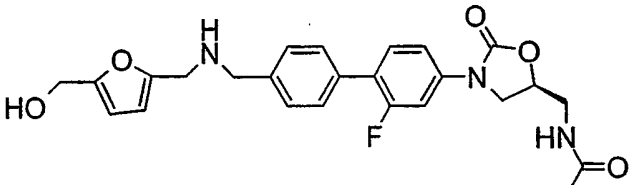
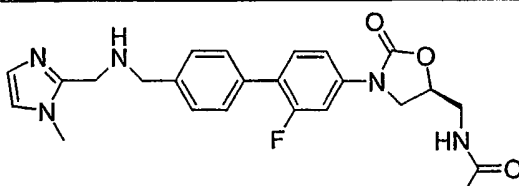
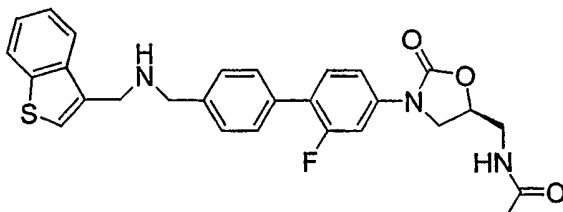
	N-(3-{2-Fluoro-4'-[(methyl-pyridin-3-ylmethyl-amino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4048	
	N-[3-(2-Fluoro-4'-{[(1H-pyrrol-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4049	
	N-[3-(2-Fluoro-4'-{[(1-methyl-1H-indol-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4050	
	1H-Indole-6-carboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-methyl-amide
4051	
	1-Methyl-1H-pyrrole-2-carboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-methyl-amide
4052	

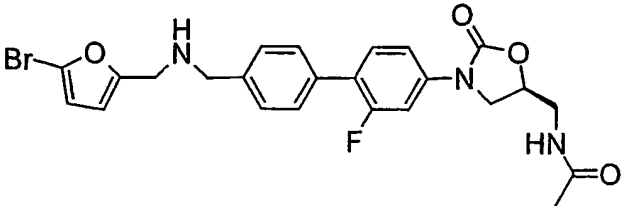
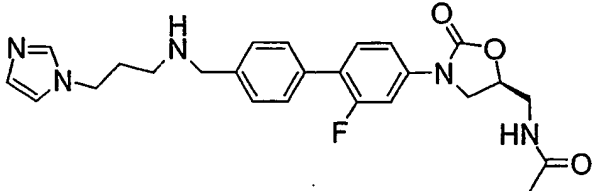
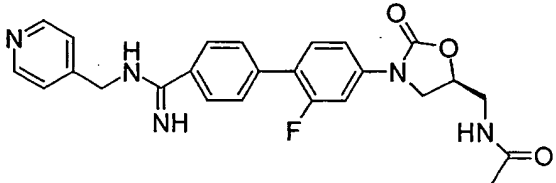
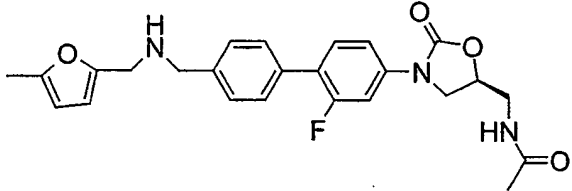
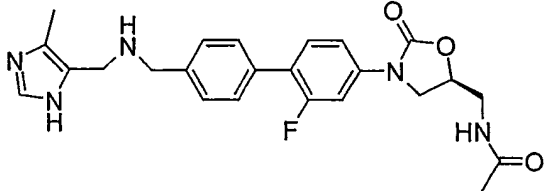
	N-{3-[3-Fluoro-4-(5-[[pyridin-4-ylmethyl]-amino]-methyl)-pyridin-2-yl]-phenyl}-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4053	
	N-{3-[3-Fluoro-4-(5-[[furan-3-ylmethyl]-amino]-methyl)-pyridin-2-yl]-phenyl}-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4054	
	N-[3-(2-Fluoro-4'-{[(6-methoxy-pyridin-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4055	
	N-[3-(2-Fluoro-4'-{[(6-methoxy-pyridin-3-ylmethyl)-methyl-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4056	
	N-(3-{4'-[(2,5-Bis-trifluoromethyl-benzylamino)-methyl]-2-fluoro-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4057	

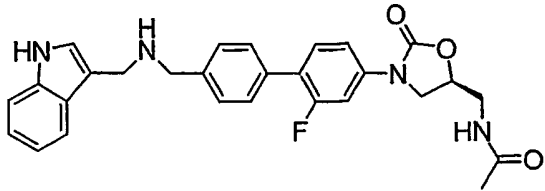
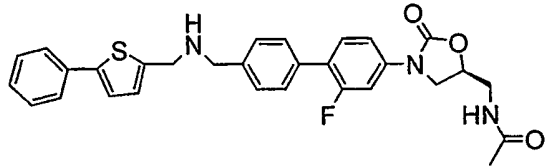
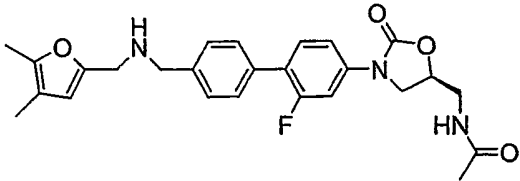
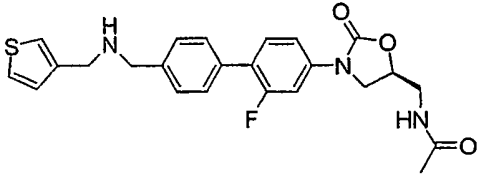
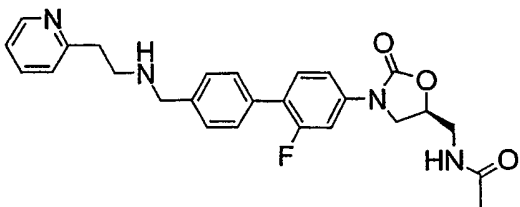
	N-[3-(2-Fluoro-4'-{[(6-methyl-2,4-dioxo-1,2,3,4-tetrahydro-pyrimidin-5-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4058	
	N-[3-(2-Fluoro-4'-{[(furan-3-ylmethyl)-amino]-methyl}-2'-methoxy-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4059	
	N-[3-(2-Fluoro-4'-{[(1-methyl-1H-pyrrol-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4060	
	N-[3-(2-Fluoro-4'-{[(isoquinolin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4061	
	N-(3-{2-Fluoro-4'-[(furan-3-ylmethyl-methyl-amino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4062	
	N-(3-{4'-[(4-Dimethylamino-benzylamino)-methyl]-2-fluoro-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide

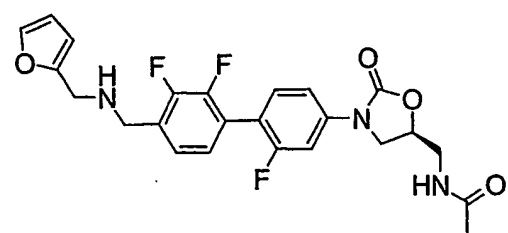
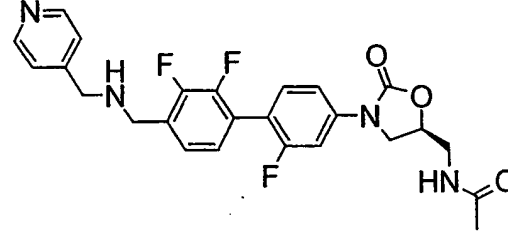
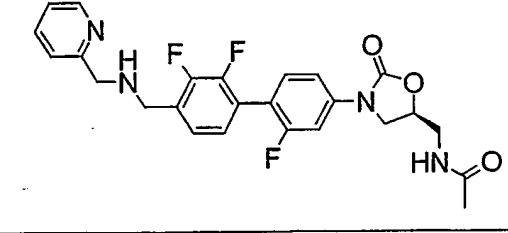
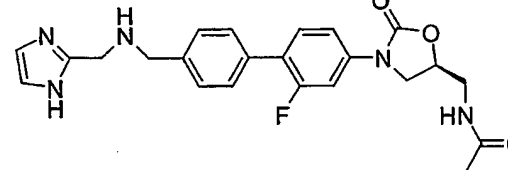
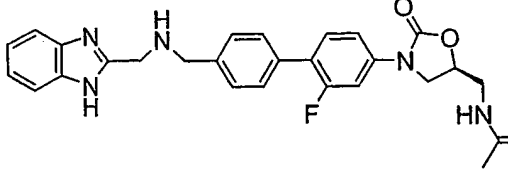
4063	
	N-(3-{4'-[(4-Chloro-benzylamino)-methyl]-2-fluoro-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4064	
	N-(3-{4'-[(2,4-Dichloro-benzylamino)-methyl]-2-fluoro-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4065	
	N-[3-(2-Fluoro-4'-{[(isoquinolin-5-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4066	
	N-[3-(2-Fluoro-4'-{[(3H-imidazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4067	
	N-[3-(2-Fluoro-4'-{[(3H-imidazol-4-ylmethyl)-methyl-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

4068	
	N-[3-(2-Fluoro-4'-{[(1H-imidazol-4-ylmethyl)-(3H-imidazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4069	
	N-[3-(2-Fluoro-4'-{[(5-nitro-furan-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4070	
	N-(3-{4'-[(3-Cyano-benzylamino)-methyl]-2-fluoro-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4071	
	N-[3-(2-Fluoro-4'-{[(quinolin-6-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4072	
	N-[3-(2-Fluoro-4'-{[(6-methyl-pyridin-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

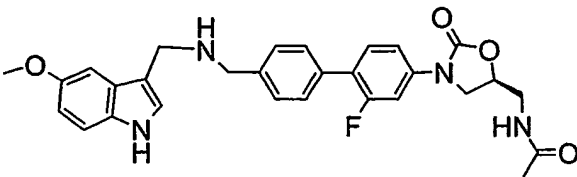
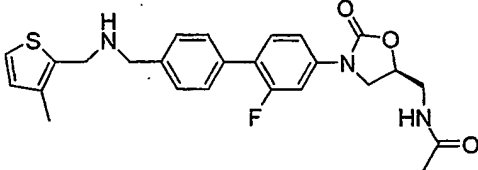
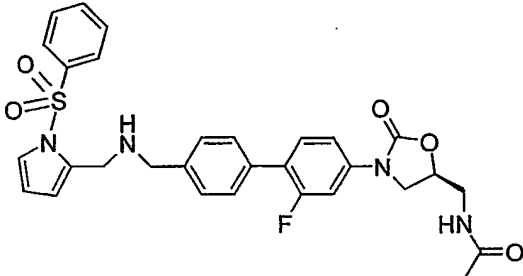
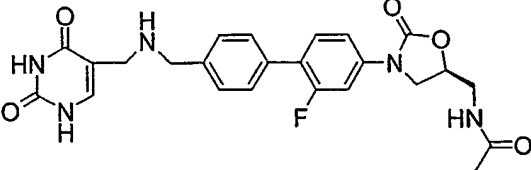
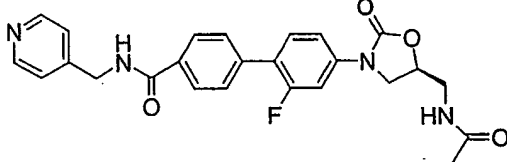
4073	
	N-[3-(3-Fluoro-4-(6-((pyridin-4-ylmethyl)-amino)-methyl)-pyridin-3-yl)-phenyl]-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4074	
	N-[3-(2-Fluoro-4'-((thiazol-2-ylmethyl)-amino)-methyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4075	
	N-[3-(2-Fluoro-4'-((5-hydroxymethyl-furan-2-ylmethyl)-amino)-methyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4076	
	N-[3-(2-Fluoro-4'-((1-methyl-1H-imidazol-2-ylmethyl)-amino)-methyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4077	
	N-[3-(4'-((Benzo[b]thiophen-3-ylmethyl)-amino)-methyl)-2-fluoro-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

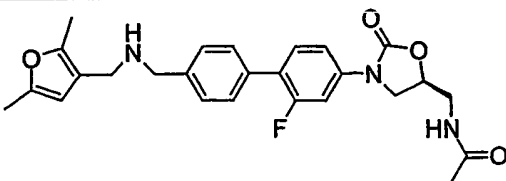
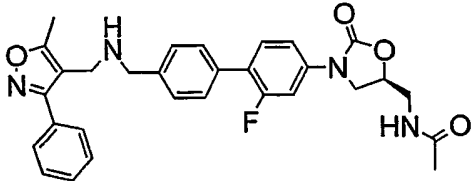
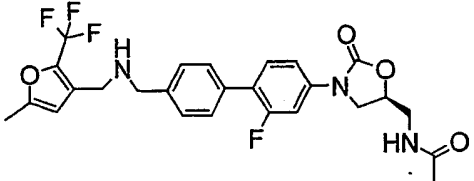
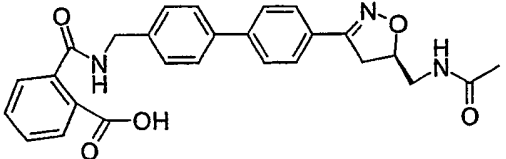
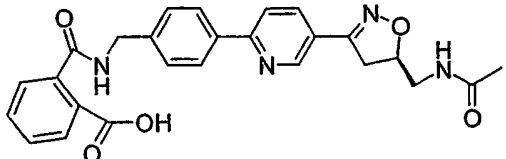
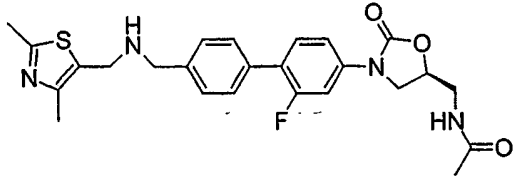
4078	
	N-[3-(4'-{[(5-Bromo-furan-2-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4079	
	N-(3-{2-Fluoro-4'-[(3-imidazol-1-yl-propylamino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4080	
	N-{3-[2-Fluoro-4'-(N-pyridin-4-ylmethyl-carbamimidoyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4081	
	N-[3-(2-Fluoro-4'-{[(5-methyl-furan-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4082	
	N-[3-(2-Fluoro-4'-{[(5-methyl-3H-imidazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

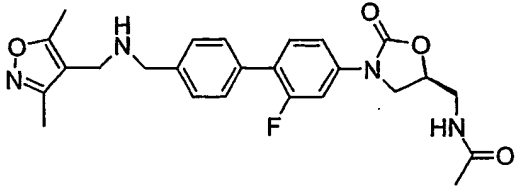
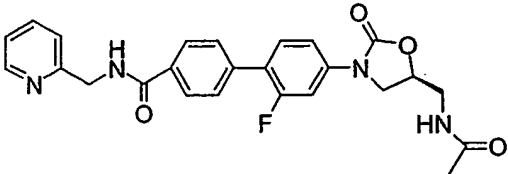
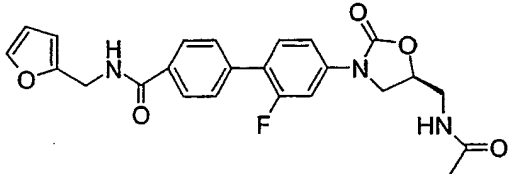
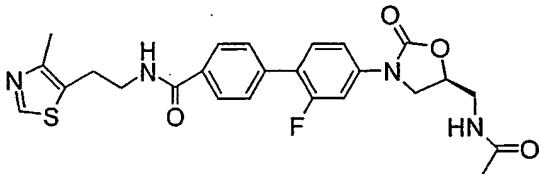
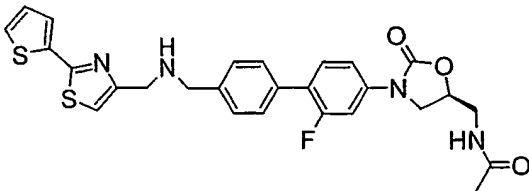
4083	
	N-[3-(2-Fluoro-4'-{[(1H-indol-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4084	
	N-[3-(2-Fluoro-4'-{[(5-phenyl-thiophen-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4085	
	N-[3-(2-Fluoro-4'-{[(4,5-Dimethyl-furan-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4086	
	N-[3-(2-Fluoro-4'-{[(thiophen-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4087	
	N-(3-{2-Fluoro-4'-[(2-pyridin-2-yl-ethylamino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide

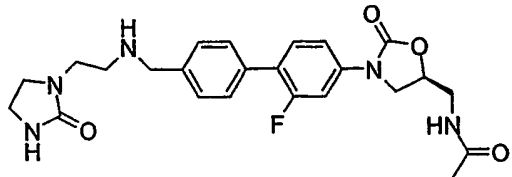
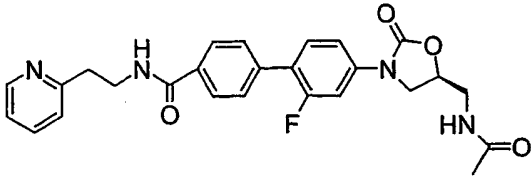
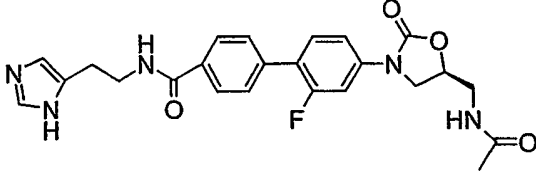
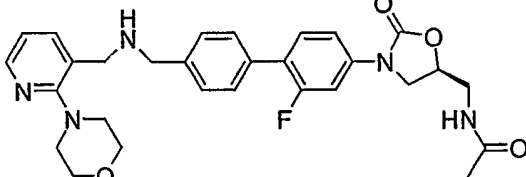
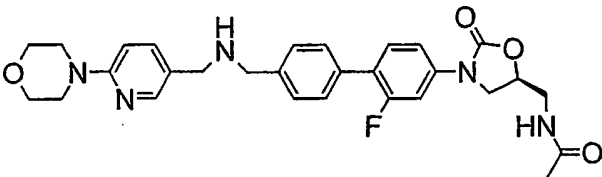
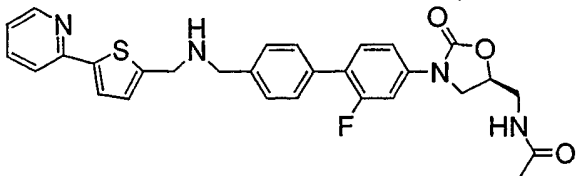
4088	
	N-[2-Oxo-3-(2,2',3'-trifluoro-4'-{[(furan-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-oxazolidin-5-(S)-ylmethyl]-acetamide
4089	
	N-[2-Oxo-3-(2,2',3'-trifluoro-4'-{[(pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-oxazolidin-5-(S)-ylmethyl]-acetamide
4090	
	N-[2-Oxo-3-(2,2',3'-trifluoro-4'-{[(pyridin-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-oxazolidin-5-(S)-ylmethyl]-acetamide
4091	
	N-[3-(2-Fluoro-4'-{[(1H-imidazol-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4092	
	N-[3-(4'-{[(1H-Benzoimidazol-2-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

4093	
	N-(3-{2-Fluoro-4'-[(4-sulfamoyl-benzylamino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4094	
	N-[3-(2-Fluoro-4'-{[2-(4-sulfamoyl-phenyl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4095	
	N-[3-(2-Fluoro-4'-{[(3-hydroxy-5-hydroxymethyl-2-methyl-pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4096	
	N-[3-(2-Fluoro-4'-{[2-(4-methyl-thiazol-5-yl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4097	
	N-{3-[2-Fluoro-4'-(N-pyridin-2-ylmethyl-carbamimidoyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide

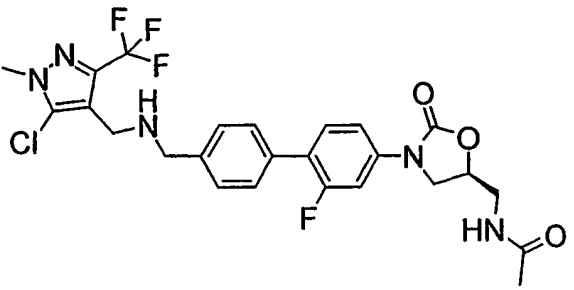
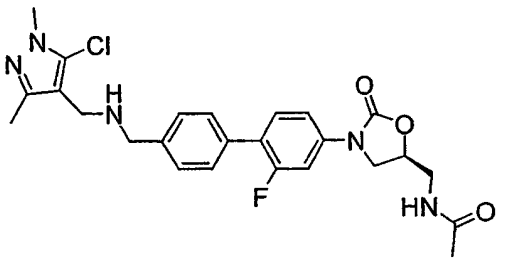
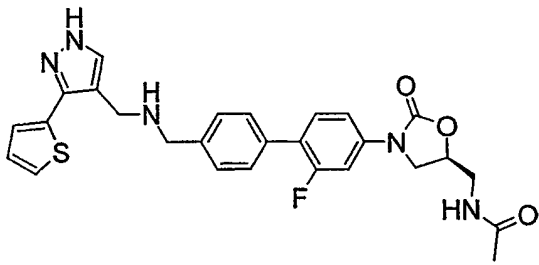
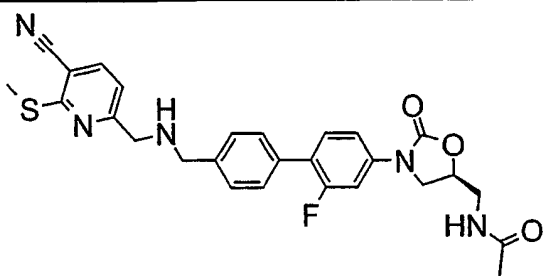
4098	
	N-[3-(2-Fluoro-4'-{[(5-methoxy-1H-indol-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4099	
	N-[3-(2-Fluoro-4'-{[(3-methyl-thiophen-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4100	
	N-[3-(4'-{[(1-Benzenesulfonyl-1H-pyrrol-2-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4101	
	N-[3-(4'-{[(2,4-Dioxo-1,2,3,4-tetrahydro-pyrimidin-5-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4102	
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-carboxylic acid (pyridin-4-ylmethyl)-amide

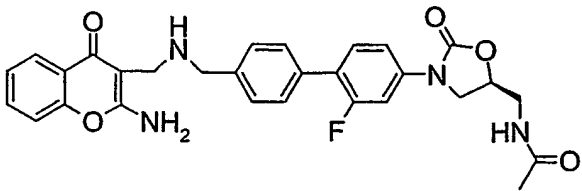
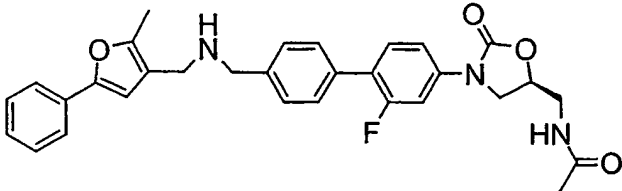
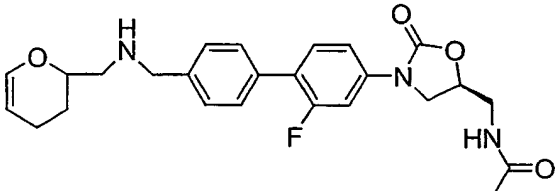
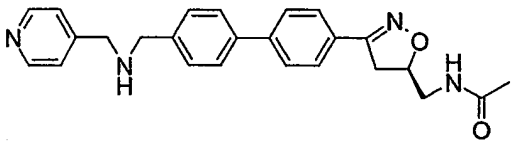
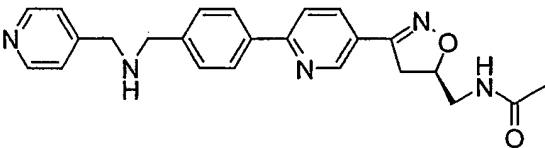
4103	
	N-[3-(4'-{[(2,5-Dimethyl-furan-3-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4104	
	N-[3-(2-Fluoro-4'-{[(5-methyl-3-phenyl-isoxazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4105	
	N-[3-(2-Fluoro-4'-{[(5-methyl-2-trifluoromethyl-furan-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4106	
	N-{4'-[5-(R)-(Acetylamino-methyl)-4,5-dihydro-isoxazol-3-yl]-biphenyl-4-ylmethyl}-phthalamic acid
4107	
	N-(4-{5-[5-(R)-(Acetylamino-methyl)-4,5-(S)-dihydro-isoxazol-3-yl]-pyridin-2-yl}-benzyl)-phthalamic acid
4108	

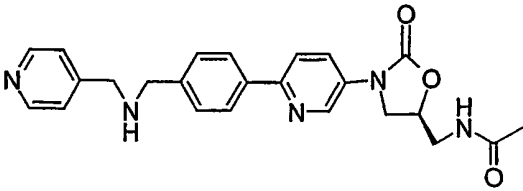
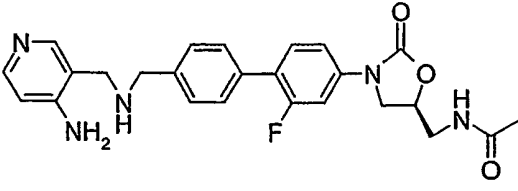
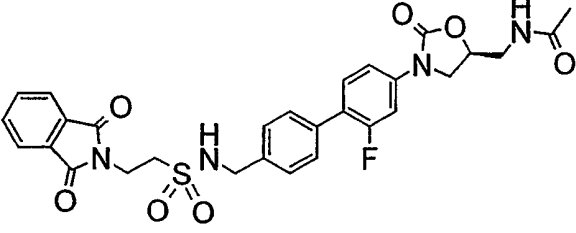
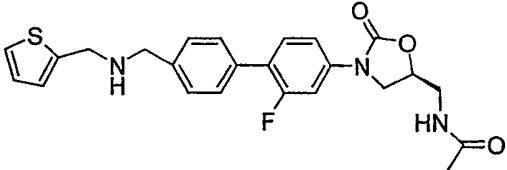
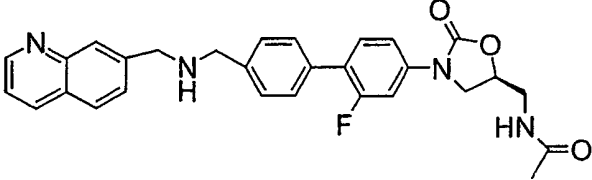
	N-[3-(4'-{[(2,4-Dimethyl-thiazol-5-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4109	
	N-[3-(4'-{[(3,5-Dimethyl-isoxazol-4-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4110	
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-carboxylic acid (pyridin-2-ylmethyl)-amide
4111	
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-carboxylic acid (furan-2-ylmethyl)-amide
4112	
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-carboxylic acid [2-(4-methyl-thiazol-5-yl)-ethyl]-amide
4113	
	N-[3-(2-Fluoro-4'-{[(2-thiophen-2-yl-thiazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

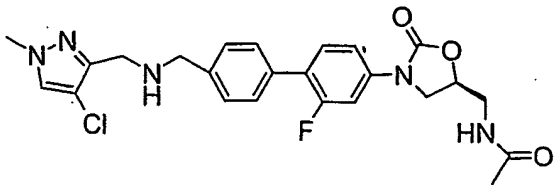
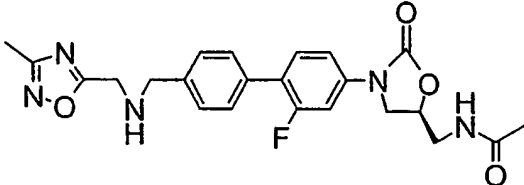
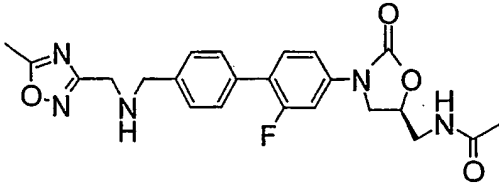
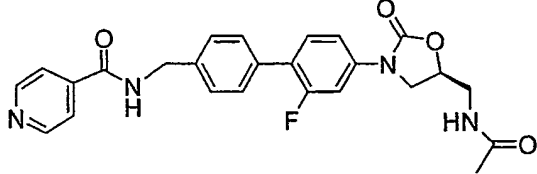
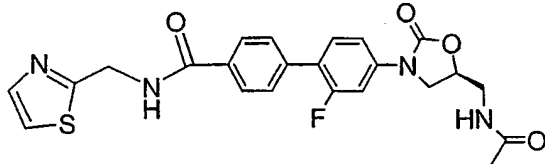
4114	
	N-[3-(2-Fluoro-4'-{[2-(2-oxo-imidazolidin-1-yl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4115	
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-carboxylic acid (2-pyridin-2-yl-ethyl)-amide
4116	
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-carboxylic acid [2-(3H-imidazol-4-yl)-ethyl]-amide
4117	
	N-[3-(2-Fluoro-4'-{[(2-morpholin-4-yl-pyridin-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4118	
	N-[3-(2-Fluoro-4'-{[(6-morpholin-4-yl-pyridin-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4119	

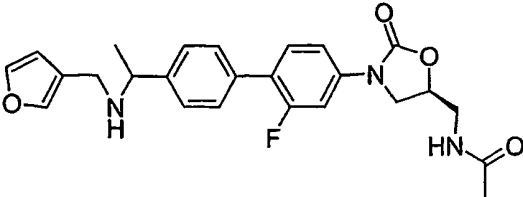
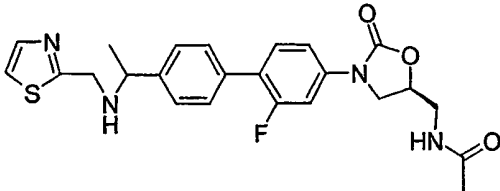
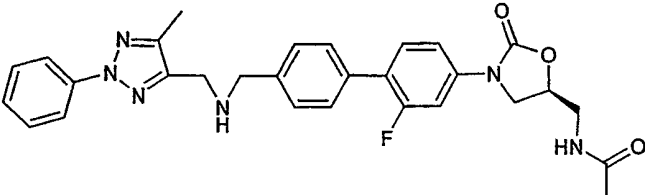
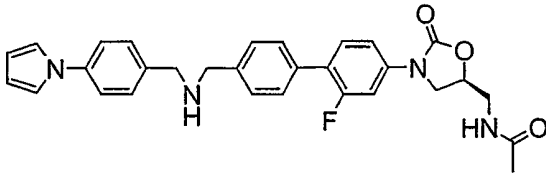
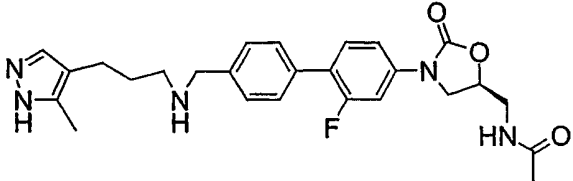
	N-[3-(2-Fluoro-4'-{[(5-pyridin-2-yl-thiophen-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4120	
	5-[(4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl)-amino)-methyl]-2-methyl-furan-3-carboxylic acid methyl ester
4121	
	N-[3-(4'-{[(Benzothiazol-2-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4122	
	N-[3-(2-Fluoro-4'-{[(2-phenyl-thiazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4123	
	N-[3-(2-Fluoro-4'-{[(2-phenyl-1H-imidazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4124	

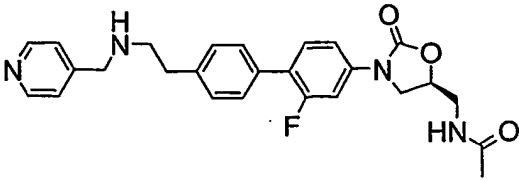
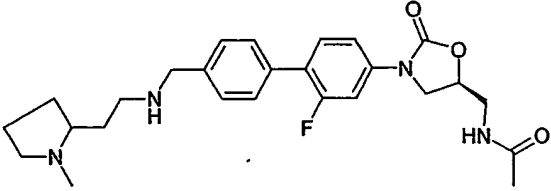
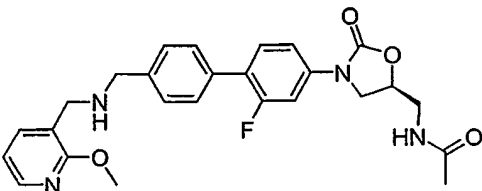
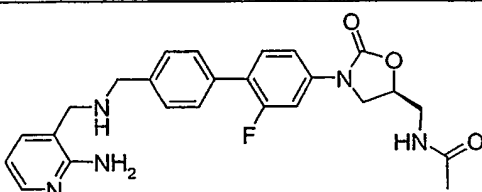
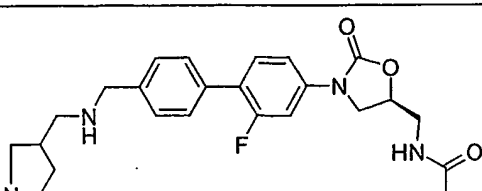
	N-[3-(4'-{[(2-Ethyl-3H-imidazol-4-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4125	
	N-[3-(4'-{[(5-Chloro-1-methyl-3-trifluoromethyl-1H-pyrazol-4-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4126	
	N-[3-(4'-{[(5-Chloro-1,3-dimethyl-1H-pyrazol-4-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4127	
	N-[3-(2-Fluoro-4'-{[(3-thiophen-2-yl-1H-pyrazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4128	

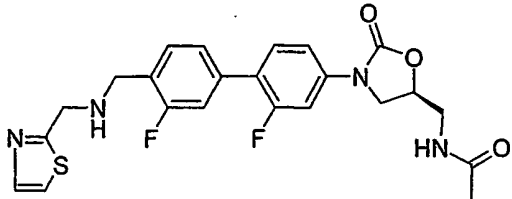
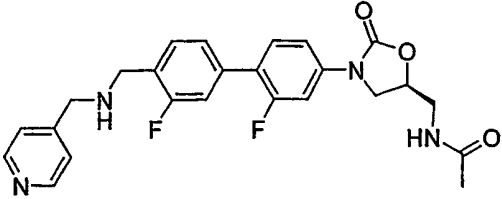
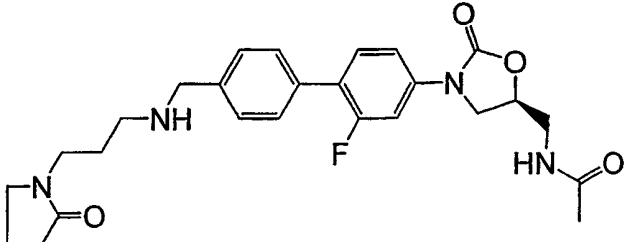
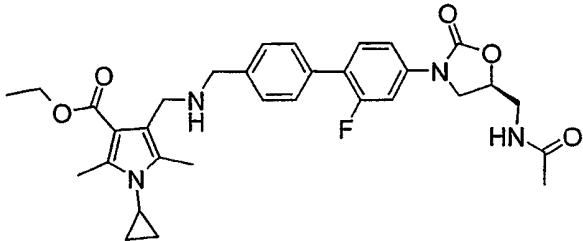
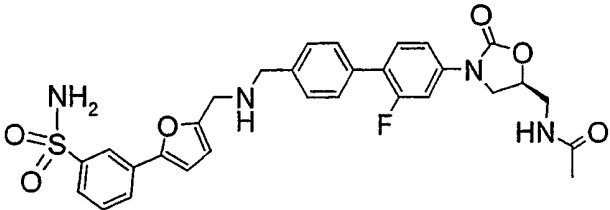
	N-[3-(4'-{[(5-Cyano-6-methylsulfanyl-pyridin-2-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4129	
	N-[3-(4'-{[(2-Amino-4-oxo-4H-chromen-3-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4130	
	N-[3-(2-Fluoro-4'-{[(2-methyl-5-phenyl-furan-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4131	
	N-[3-(4'-{[(3,4-Dihydro-2H-pyran-2-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4132	
	N-[3-(4'-{[(Pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-4,5-dihydro-isoxazol-5-(R)-ylmethyl]-acetamide
4133	
	N-{3-[6-(4-{[(Pyridin-4-ylmethyl)-amino]-methyl}-phenyl)-pyridin-3-yl]-4,5-dihydro-isoxazol-5-(R)-ylmethyl}-acetamide

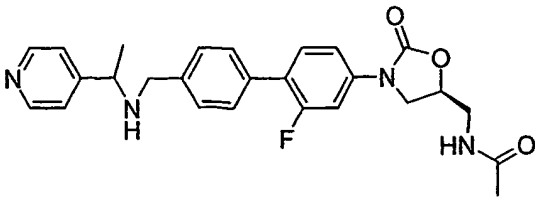
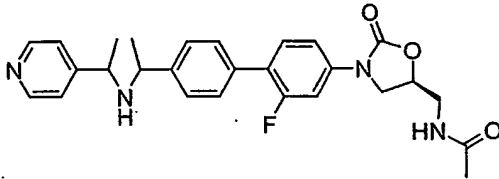
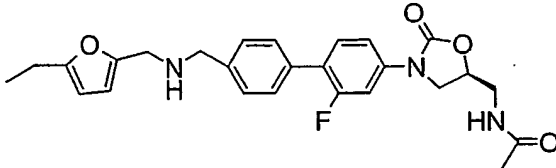
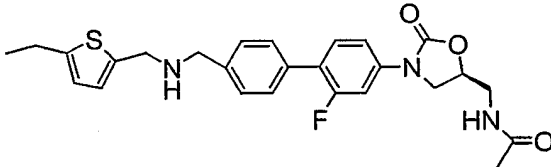
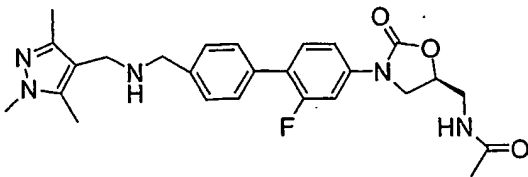
4134	
	N-[2-Oxo-3-[6-(4-[(pyridin-4-ylmethyl)-amino]-methyl)-phenyl]-pyridin-3-yl]-oxazolidin-5-(S)-ylmethyl]-acetamide
4135	
	N-[3-(4'-{[(4-Amino-pyridin-3-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4136	
	N-[3-(4'-{[2-(1,3-Dioxo-1,3-dihydro-isindol-2-yl)-ethanesulfonylamino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4137	
	N-[3-(2-Fluoro-4'-{[(thiophen-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4138	
	N-[3-(2-Fluoro-4'-{[(quinolin-7-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

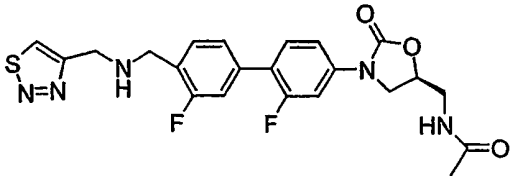
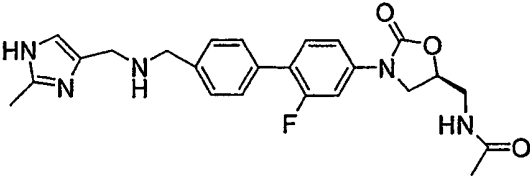
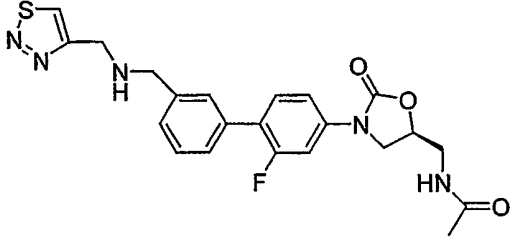
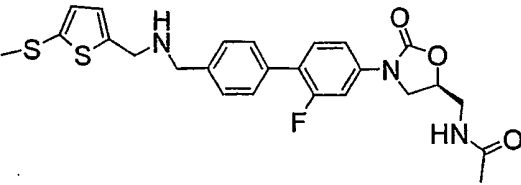
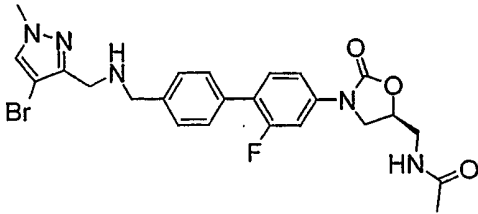
4139	
	N-[3-(4'-{[(4-Chloro-1-methyl-1H-pyrazol-3-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4140	
	N-[3-(2-Fluoro-4'-{[(3-methyl-[1,2,4]oxadiazol-5-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4141	
	N-[3-(2-Fluoro-4'-{[(5-methyl-[1,2,4]oxadiazol-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4142	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-isonicotinamide
4143	
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-carboxylic acid (thiazol-2-ylmethyl)-amide

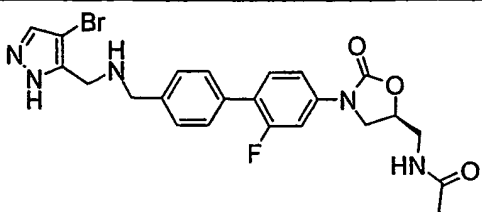
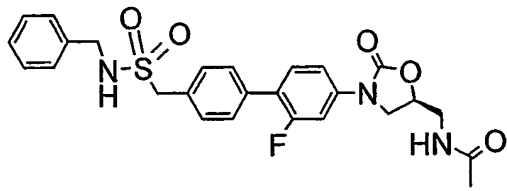
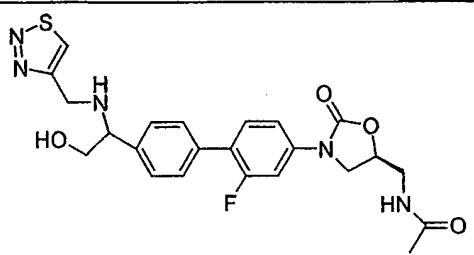
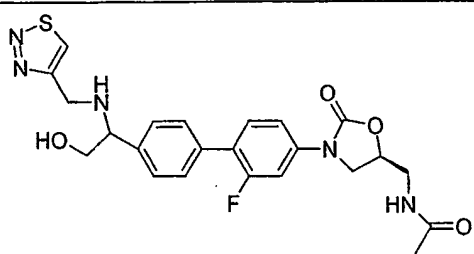
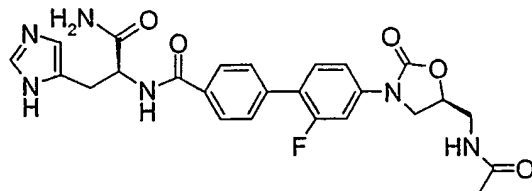
4144	
	N-[3-(2-Fluoro-4'-{1-(R/S)-[(furan-3-ylmethyl)-amino]-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4145	
	N-[3-(2-Fluoro-4'-{1-(R/S)-[(thiazol-2-ylmethyl)-amino]-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4146	
	N-[3-(2-Fluoro-4'-{1-(R/S)-[(5-methyl-2-phenyl-2H-[1,2,3]triazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4147	
	N-[3-(2-Fluoro-4'-{1-(R/S)-[(4-pyrrol-1-yl-benzylamino)-methyl]-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4148	
	N-[3-(2-Fluoro-4'-{1-(R/S)-[(3-(5-methyl-1H-pyrazol-4-yl)-propylamino)-methyl]-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

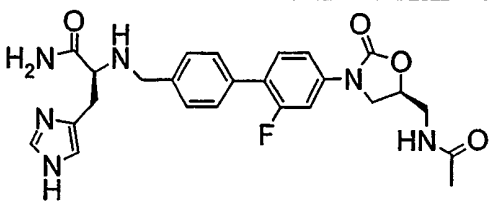
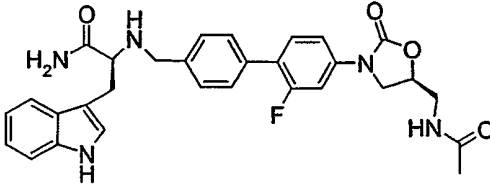
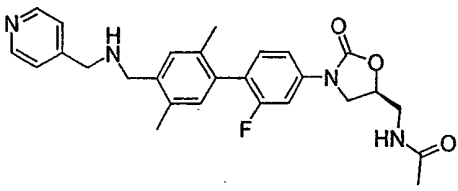
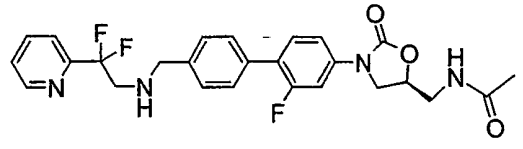
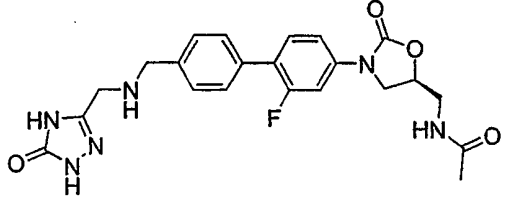
4149	
	N-[3-(2-Fluoro-4'-{2-[(pyridin-4-ylmethyl)-amino]-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4150	
	N-[3-(2-Fluoro-4'-{2-(R/S)-(1-methyl-pyrrolidin-2-yl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4151	
	N-[3-(2-Fluoro-4'-{[(2-methoxy-pyridin-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4152	
	N-[3-(4'-{[(2-Amino-pyridin-3-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4153	
	N-[3-(2-Fluoro-4'-{[(pyrrolidin-3-(R/S)-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

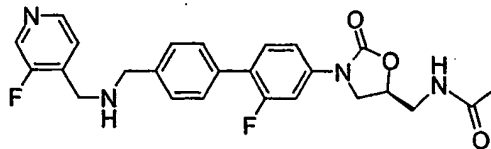
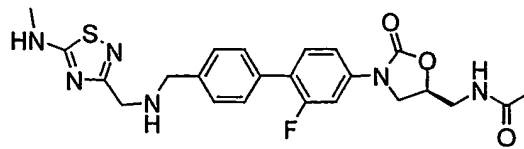
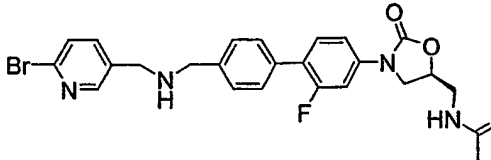
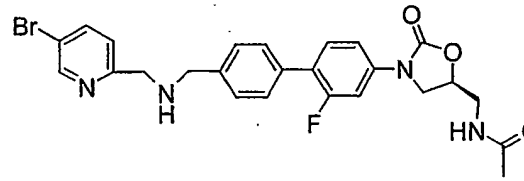
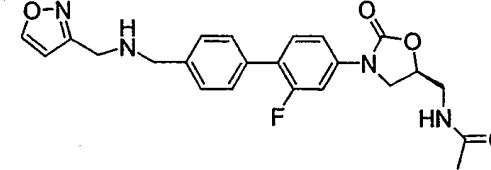
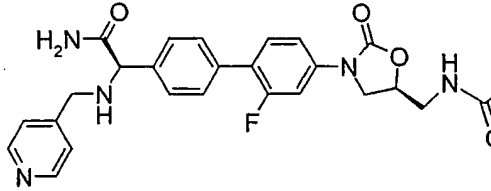
4154	
	N-[3-(2,3'-Difluoro-4'-{[(thiazol-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4155	
	N-[3-(2,3'-Difluoro-4'-{[(pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4156	
	N-[3-(2-Fluoro-4'-{[3-(2-oxo-pyrrolidin-1-yl)-propylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4157	
	4-[(4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl)-amino]-methyl-1-cyclopropyl-2,5-dimethyl-1H-pyrrole-3-carboxylic acid ethyl ester
4158	

	N-{3-[2-Fluoro-4'-{[(5-(3-sulfamoyl-phenyl)-furan-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4159	
	N-(3-{2-Fluoro-4'-[(1-pyridin-4-(R/S)-yl-ethylamino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4160	
	N-(3-{2-Fluoro-4'-[1-(R/S)-(1-pyridin-4-(R/S)-yl-ethylamino)-ethyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4161	
	N-[3-(4'-{[(5-Ethyl-furan-2-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4162	
	N-[3-(4'-{[(5-Ethyl-thiophen-2-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4163	
	N-[3-(2-Fluoro-4'-{[(1,3,5-trimethyl-1H-pyrazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

4164	
	N-[3-(2,3'-Difluoro-4'-{[(1,2,3]thiadiazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4165	
	N-[3-(2-Fluoro-4'-{[(2-methyl-1H-imidazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4166	
	N-[3-(2-Fluoro-3'-{[(1,2,3]thiadiazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4167	
	N-[3-(2-Fluoro-4'-{[(5-methylsulfanyl-thiophen-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4168	
	N-[3-(4'-{[(4-Bromo-1-methyl-1H-pyrazol-3-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

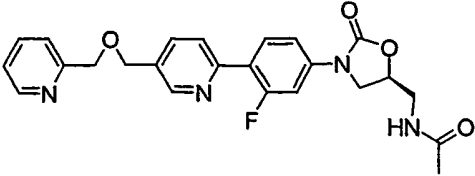
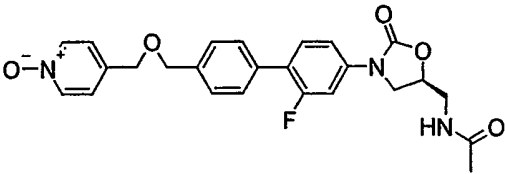
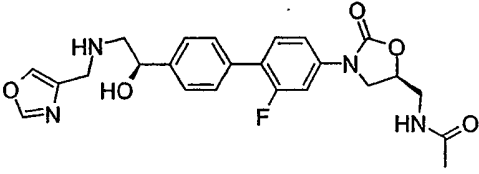
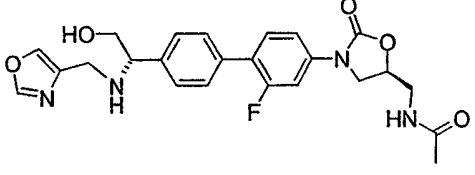
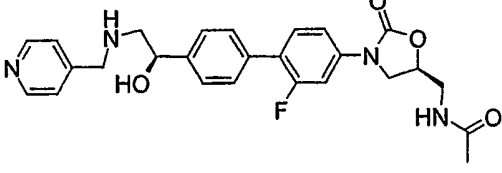
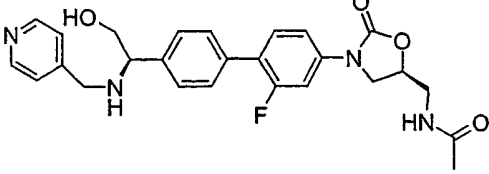
4169	
	N-[3-(4'-{[(4-Bromo-2H-pyrazol-3-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4170	
	N-{3-[4'-(Benzylsulfamoyl-methyl)-2-fluoro-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4171	
	N-[3-(2-Fluoro-4'-{2-hydroxy-1-[(1,2,3]thiadiazol-4-(R/S)-ylmethyl)-amino]-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4172	
	N-[3-(2-Fluoro-4'-{2-hydroxy-1-[(1,2,3]thiadiazol-4-(R/S)-ylmethyl)-amino]-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4173	

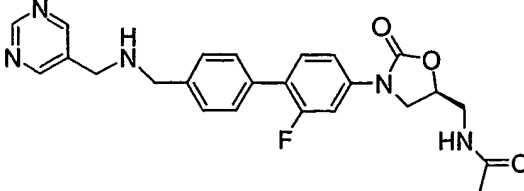
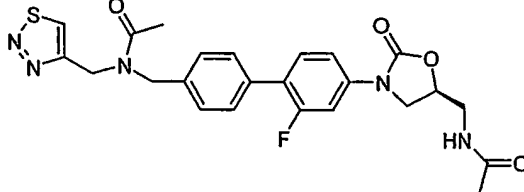
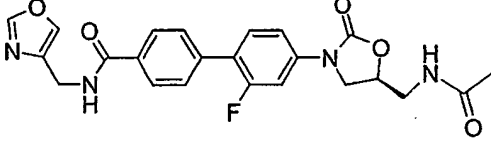
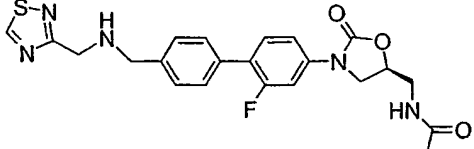
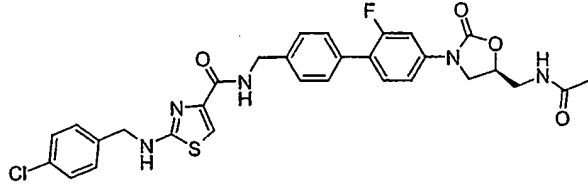
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-carboxylic acid [1-carbamoyl-2-(S)-(3H-imidazol-4-yl)-ethyl]-amide
4174	
	2-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amino)-3-(S)-(1H-imidazol-4-yl)-propionamide
4175	
	2-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amino)-3-(S)-(1H-indol-3-yl)-propionamide
4176	
	N-[3-(2-Fluoro-2',5'-dimethyl-4'-{[(pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4177	
	N-(3-{4'-[(2,2-Difluoro-2-pyridin-2-yl-ethylamino)-methyl]-2-fluoro-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4178	
	N-[3-(2-Fluoro-4'-{[(5-(S)-oxo-4,5-dihydro-1H-[1,2,4]triazol-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

4179	
	N-[3-(2-Fluoro-4'-{[(3-fluoro-pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4180	
	N-[3-(2-Fluoro-4'-{[(5-methylamino-[1,2,4]thiadiazol-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4181	
	N-[3-(4'-{[(6-Bromo-pyridin-3-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4182	
	N-[3-(4'-{[(5-Bromo-pyridin-2-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4183	
	N-[3-(2-Fluoro-4'-{[(isoxazol-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4184	
	2-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-yl}-2-(R)-[(pyridin-4-ylmethyl)-amino]-acetamide

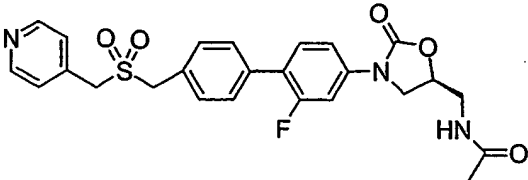
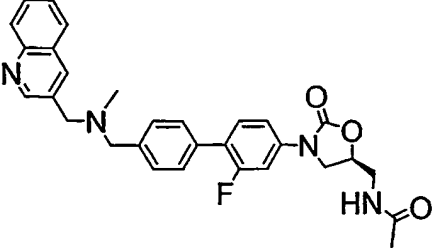
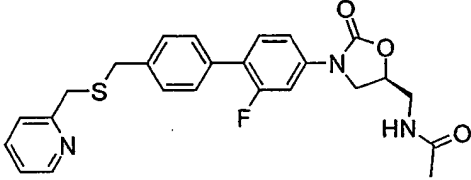
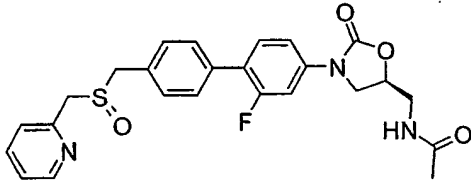
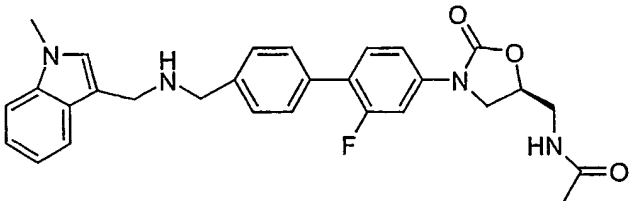
4185	
	2-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-yl}-2-(R)-[(pyridin-2-ylmethyl)-amino]-acetamide
4186	
	N-[3-(2-Fluoro-4'-{[(piperidin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4187	
	5-{4-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2-fluoro-phenyl}-pyridine-2-carboxylic acid (pyridin-2-ylmethyl)-amide
4188	
	5-{4-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2-fluoro-phenyl}-pyridine-2-carboxylic acid (pyridin-4-ylmethyl)-amide
4189	
	5-{4-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2-fluoro-phenyl}-pyridine-2-carboxylic acid (thiazol-2-ylmethyl)-amide
4190	
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-3,2'-difluoro-biphenyl-4-carboxylic acid (pyridin-2-ylmethyl)-amide
4191	

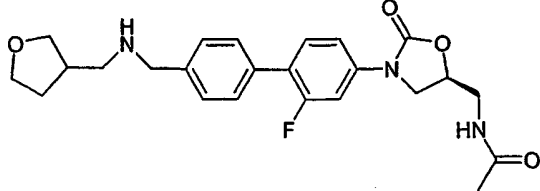
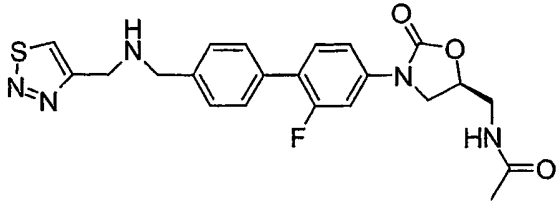
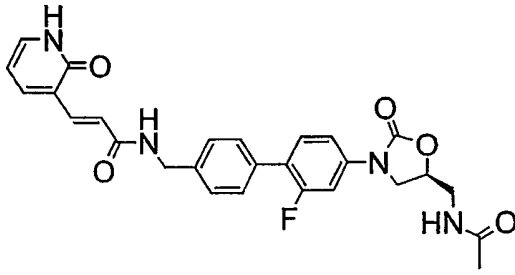
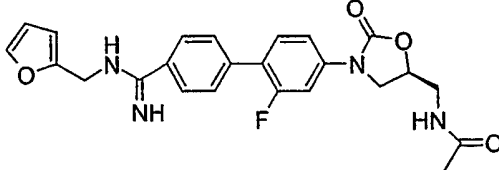
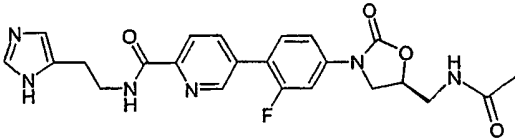
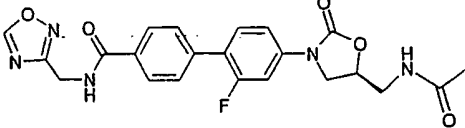
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-3,2'-difluoro-biphenyl-4-carboxylic acid [2-(3H-imidazol-4-yl)-ethyl]-amide
4192	
	N-{3-[2-Fluoro-4'-(pyridin-2-ylmethoxymethyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4193	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-3-(1-methyl-6-oxo-1,6-dihydro-pyridin-3-yl)-acrylamide
4194	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-3-(1-methyl-2-oxo-1,2-dihydro-pyridin-3-yl)-acrylamide
4195	
	N-(3-{3-Fluoro-4-[6-(pyridin-2-ylmethoxymethyl)-pyridin-3-yl]-phenyl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4196	
	N-{3-[2-Fluoro-4'-(pyridin-4-ylmethoxymethyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide

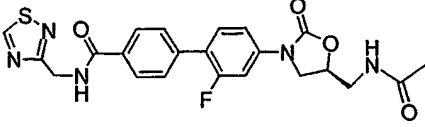
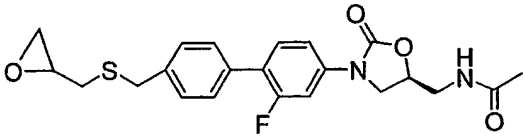
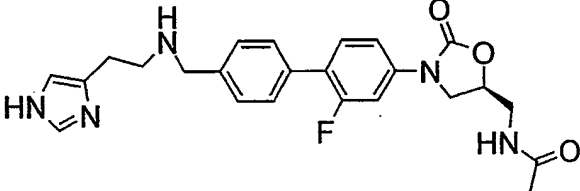
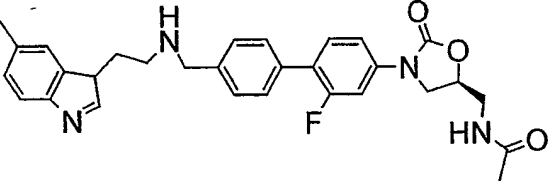
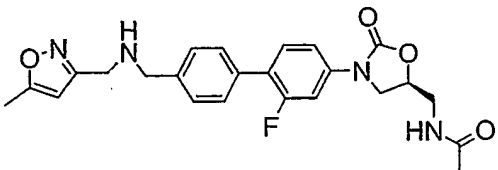
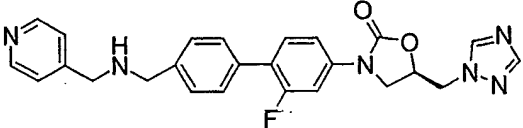
4197	
	N-(3-{3-Fluoro-4-[5-(pyridin-2-ylmethoxymethyl)-pyridin-2-yl]-phenyl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4198	
	N-{3-[2-Fluoro-4'-(1-oxy-pyridin-4-ylmethoxymethyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4199	
	N-[3-(2-Fluoro-4'-{1-(R)-hydroxy-2-[(oxazol-4-ylmethyl)-amino]-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4200	
	N-[3-(2-Fluoro-4'-{2-hydroxy-1-(S)-[(oxazol-4-ylmethyl)-amino]-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4201	
	N-[3-(2-Fluoro-4'-{1-(R)-hydroxy-2-[(pyridin-4-ylmethyl)-amino]-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4202	

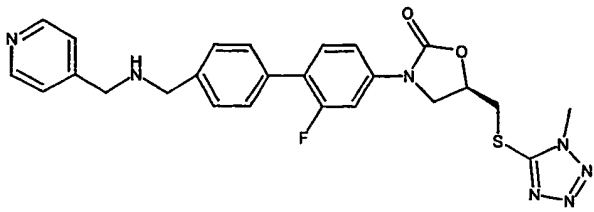
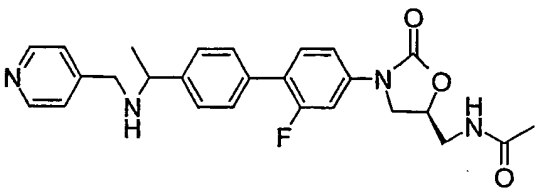
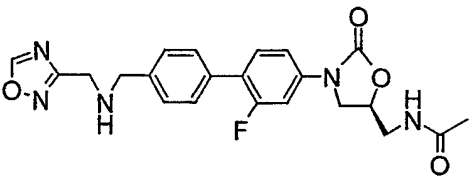
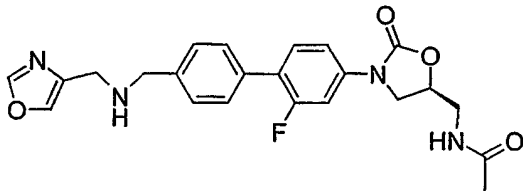
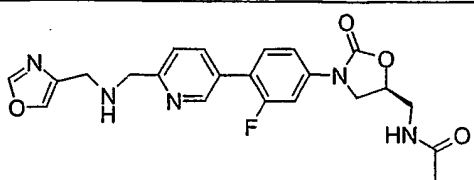
	N-[3-(2-Fluoro-4'-{2-hydroxy-1-(R)-[(pyridin-4-ylmethyl)-amino]-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4203	
	N-[3-(2-Fluoro-4'-{[(pyrimidin-5-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4204	
	N-(3-{4'-[(Acetyl-[1,2,3]thiadiazol-4-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4205	
	4'-[5-(S)-(Acetyl-amino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-carboxylic acid (oxazol-4-ylmethyl)-amide
4206	
	N-[3-(2-Fluoro-4'-{[(1,2,4]thiadiazol-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4207	
	2-(4-Chloro-benzylamino)-thiazole-4-carboxylic acid {4'-[5-(S)-(acetyl-amino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide

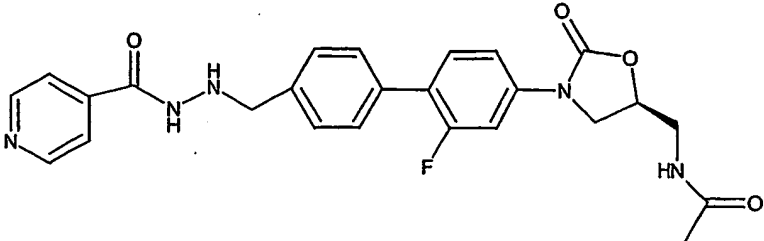
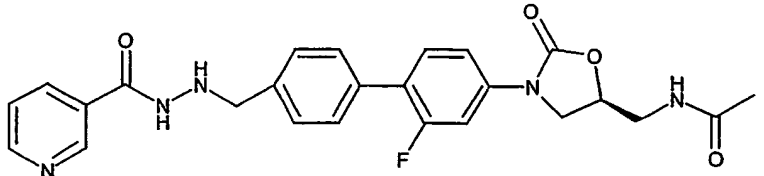
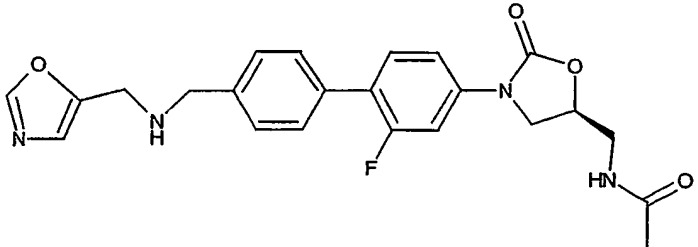
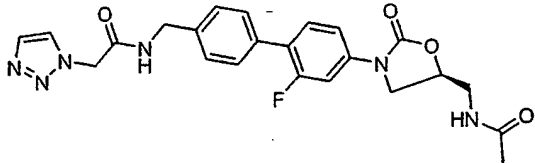
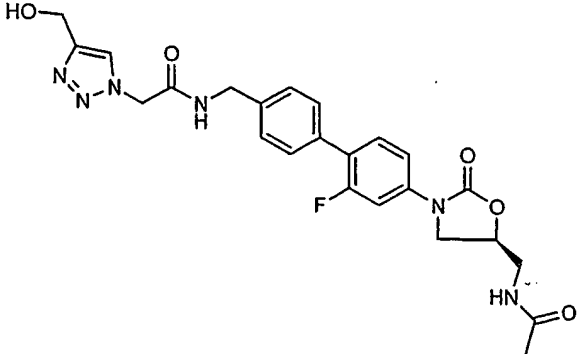
4208	
	N-[3-(2-Fluoro-4'-{[(oxazol-5-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4209	
	N-[3-(4'-{[(1,3]Dioxolan-2-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4210	
	N-(3-{2-Fluoro-4'-[(oxiranylmethyl-amino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4211	
	N-{3-[2-Fluoro-4'-(pyridin-4-ylmethylsulfanylmethyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4212	
	N-{3-[2-Fluoro-4'-(pyridin-4-ylmethanesulfinylmethyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4213	
	3-(2-Fluoro-4'-{[(pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-5-(R)-hydroxymethyl-oxazolidin-2-one

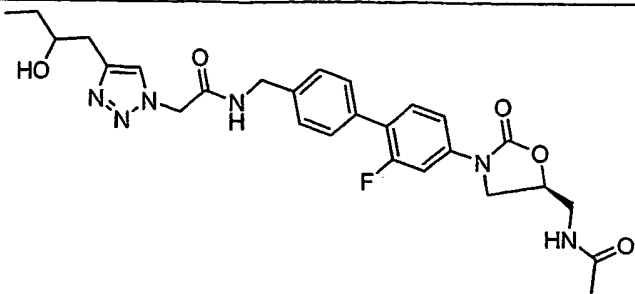
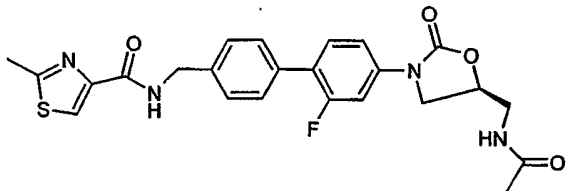
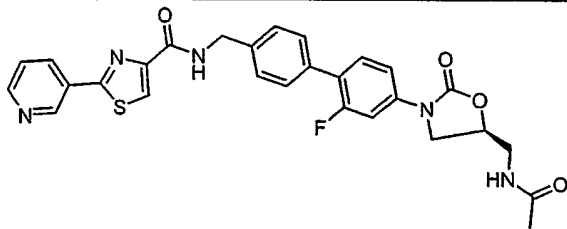
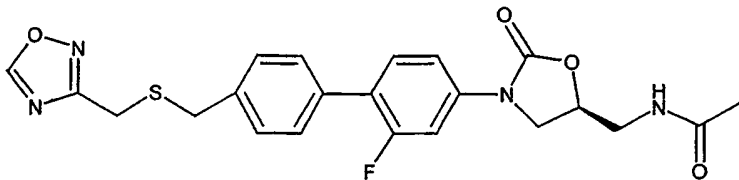
4214	
	N-{3-[2-Fluoro-4'-(pyridin-4-ylmethanesulfonylmethyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4215	
	N-(3-{2-Fluoro-4'-[(methyl-quinolin-3-ylmethyl-amino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4216	
	N-{3-[2-Fluoro-4'-(pyridin-2-ylmethanethiylmethyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4217	
	N-{3-[2-Fluoro-4'-(pyridin-2-ylmethanesulfinylmethyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4218	
	N-[3-(2-Fluoro-4'-{[(1-methyl-1H-indol-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

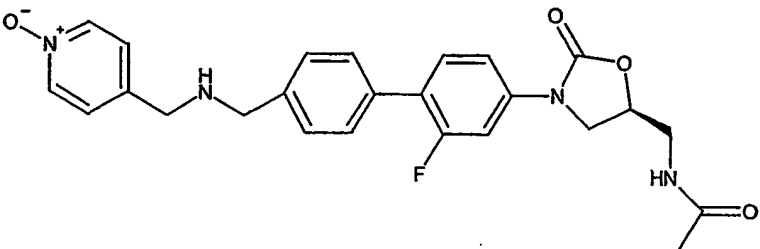
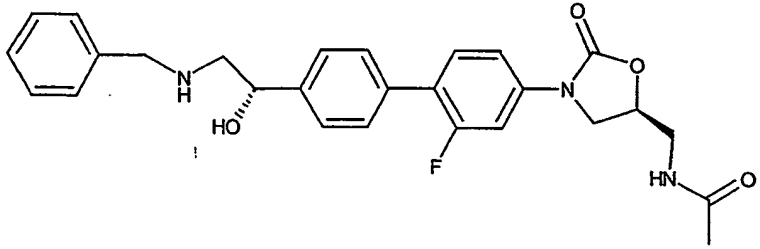
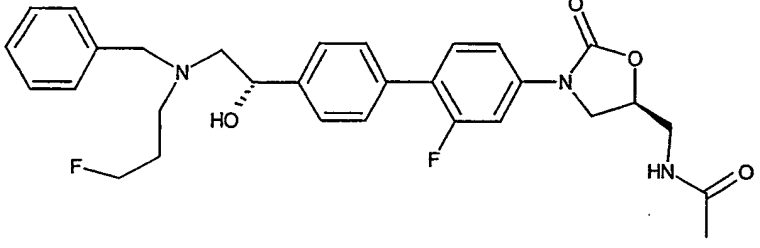
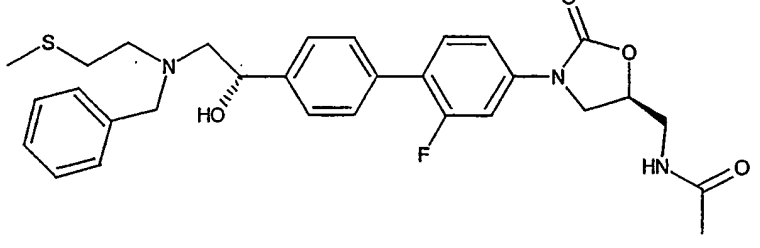
4219	
	N-[3-(2-Fluoro-4'-{[(tetrahydro-furan-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4220	
	N-[3-(2-Fluoro-4'-{[(1,2,4-triazol-5-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4221	
	N-[3-(2-Fluoro-4'-{[(thiophen-2-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4222	
	N-[3-(2-Fluoro-4'-{[(N-furan-2-ylmethyl-carbamimidoyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4223	
	5-{4-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2-fluorophenyl}-pyridine-2-carboxylic acid [2-(3H-imidazol-4-yl)-ethyl]-amide
4224	

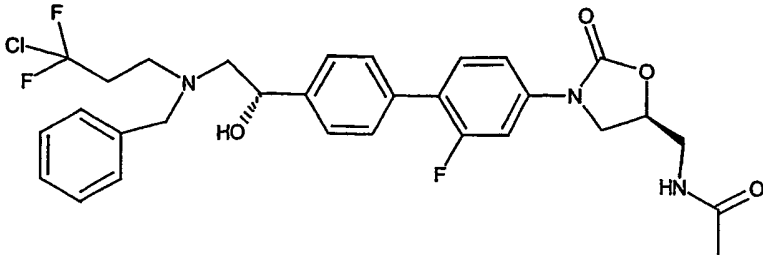
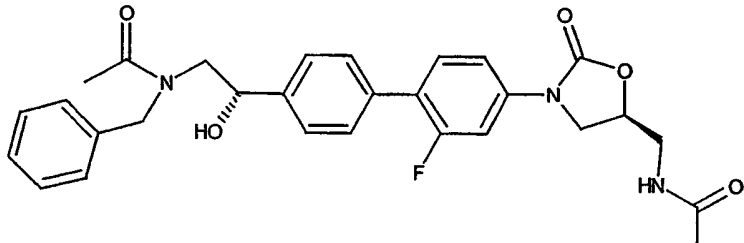
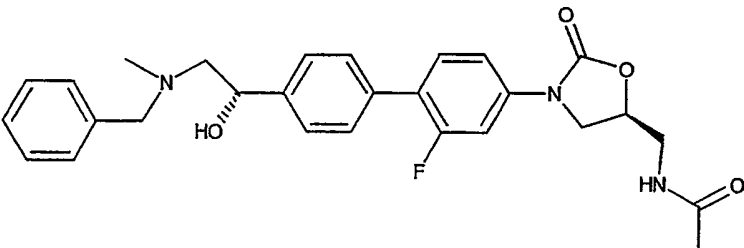
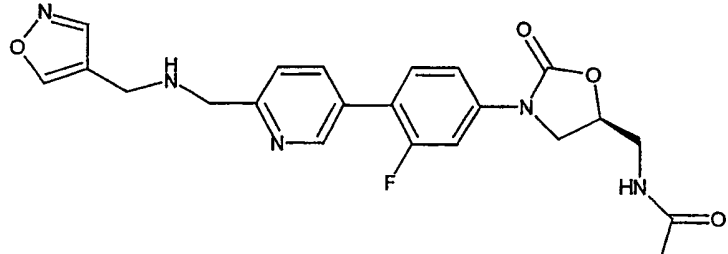
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-carboxylic acid ([1,2,4]oxadiazol-3-ylmethyl)-amide
4225	
	4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-carboxylic acid ([1,2,4]thiadiazol-3-ylmethyl)-amide
4226	
	N-[3-(2-Fluoro-4'-oxiranylmethylsulfanylmethyl-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4227	
	N-[3-(2-Fluoro-4'-{[2-(1H-imidazol-4-yl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4228	
	N-[3-(2-Fluoro-4'-{[2-(5-methyl-3H-indol-3-yl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4229	
	N-[3-(2-Fluoro-4'-{[(5-methyl-isoxazol-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4230	

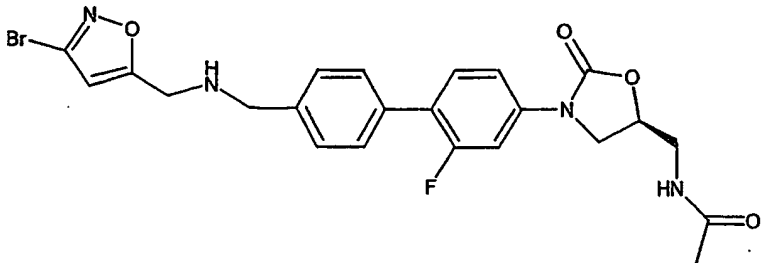
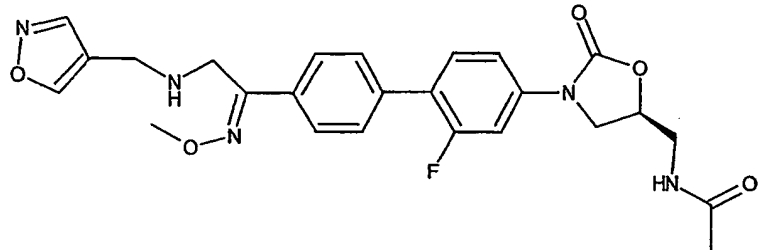
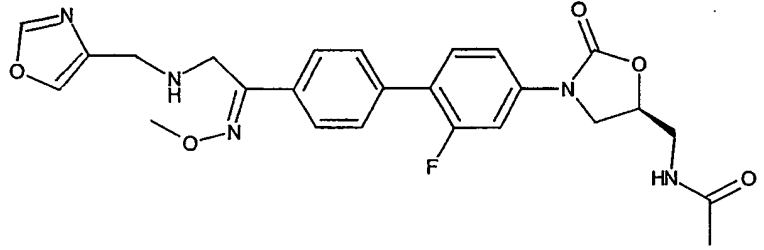
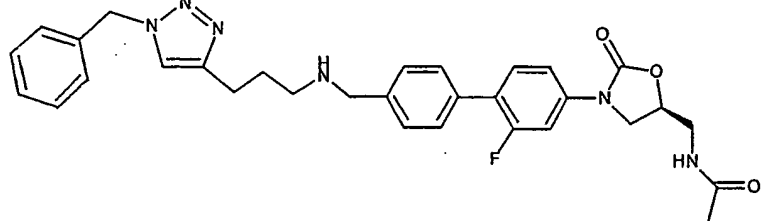
	3-(2-Fluoro-4'-{[(pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-5-(R)-[1,2,4]triazol-1-ylmethyl-oxazolidin-2-one
4231	
	3-(2-Fluoro-4'-{[(pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-5-(R)-(1-methyl-1H-tetrazol-5-ylsulfanylmethyl)-oxazolidin-2-one
4232	
	N-[3-(2-Fluoro-4'-{1-(R/S)-[(pyridin-4-ylmethyl)-amino]-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4233	
	N-[3-(2-Fluoro-4'-{[(1,2,4]oxadiazol-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4234	
	N-[3-(2-Fluoro-4'-{[(oxazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4235	
	N-{3-[3-Fluoro-4-(6-{[(oxazol-4-ylmethyl)-amino]-methyl}-pyridin-3-yl)-phenyl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide

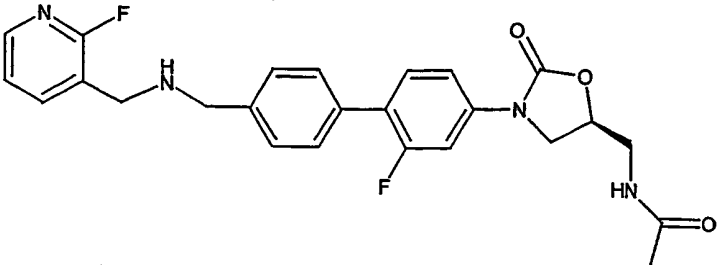
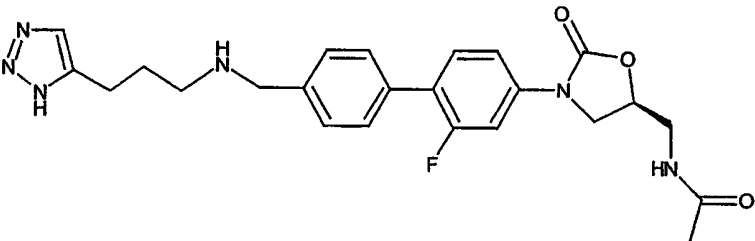
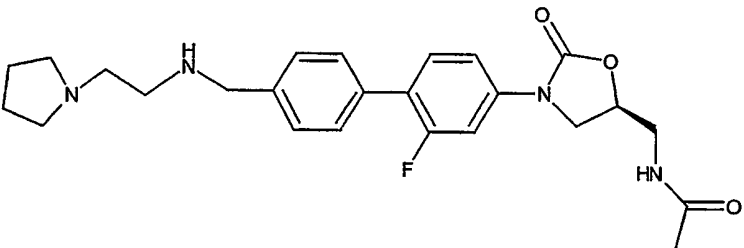
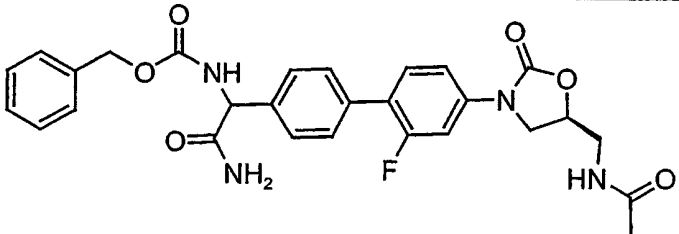
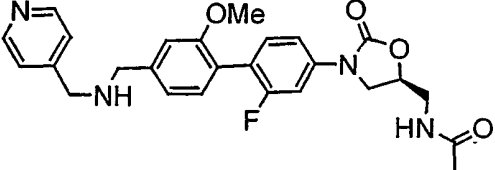
4236	
	N-(3-{2-Fluoro-4'-[N'-(pyridine-4-carbonyl)-hydrazinomethyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4237	
	N-(3-{2-Fluoro-4'-[N'-(pyridine-3-carbonyl)-hydrazinomethyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4238	
	N-[3-(2-Fluoro-4'-{[(oxazol-5-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4239	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-2-[1,2,3]triazol-1-yl-acetamide
4240	

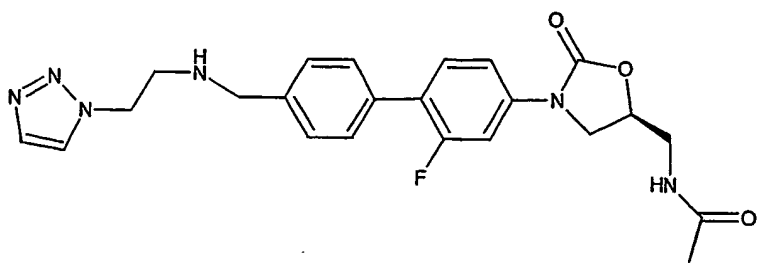
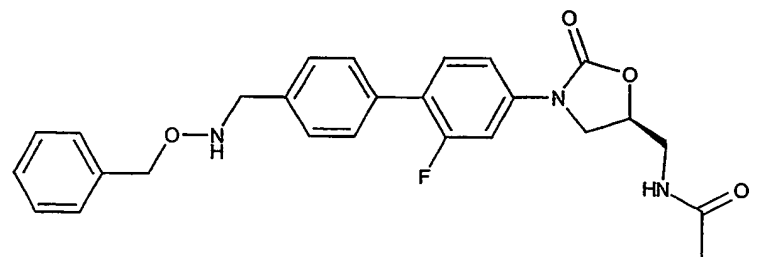
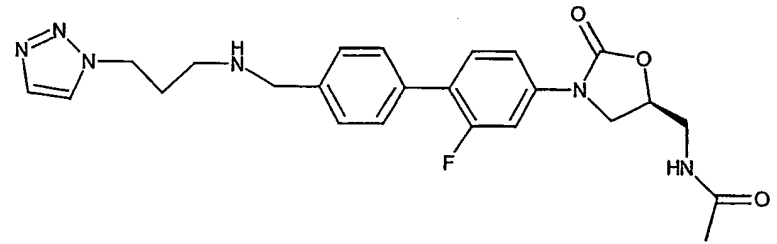
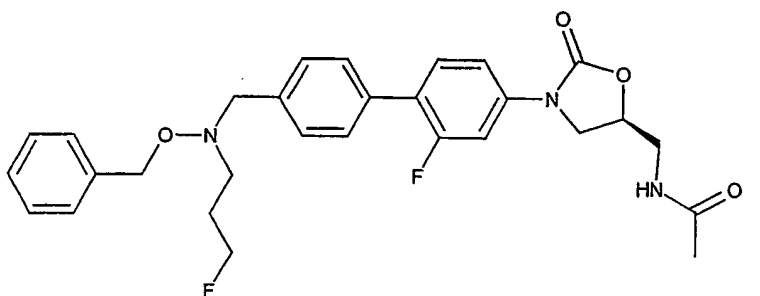
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-2-(4-hydroxymethyl-[1,2,3]triazol-1-yl)-acetamide
4241	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-2-[4-(2-hydroxy-butyl)-[1,2,3]triazol-1-yl]-acetamide
4242	
	2-Methyl-thiazole-4-carboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide
4243	
	2-Methyl-thiazole-4-carboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide
4244	
	N-{3-[2-Fluoro-4'-([1,2,4]oxadiazol-3-ylmethylsulfanylmethyl)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide

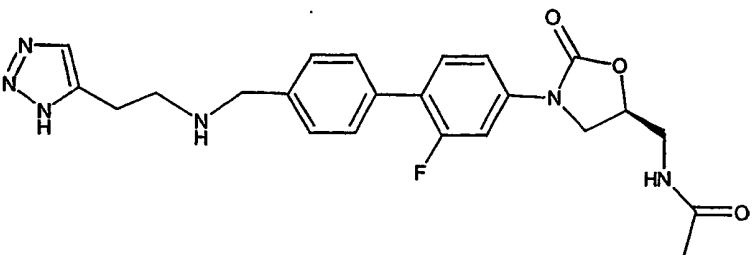
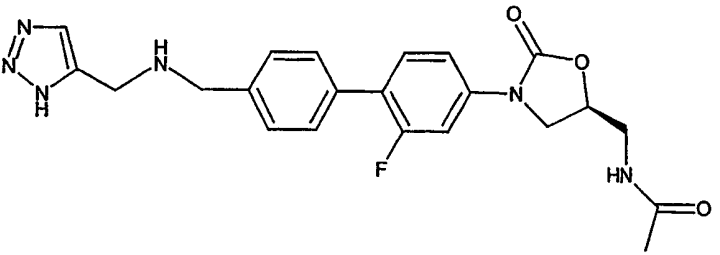
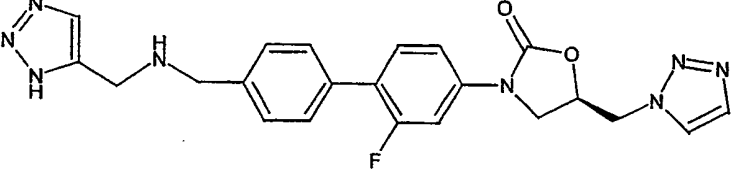
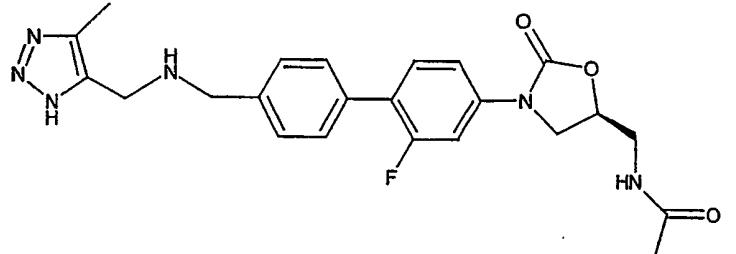
4245	
	N-[3-(2-Fluoro-4'-{[(1-oxy-pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4246	
	N-[3-(4'-(2-Benzylamino-1-(S)-hydroxy-ethyl)-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4247	
	N-[3-(4'-(2-[Benzyl-(3-fluoro-propyl)-amino]-1-(S)-hydroxy-ethyl)-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4248	
	N-[3-(4'-(2-[Benzyl-(2-methylsulfanyl-ethyl)-amino]-1-(S)-hydroxy-ethyl)-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

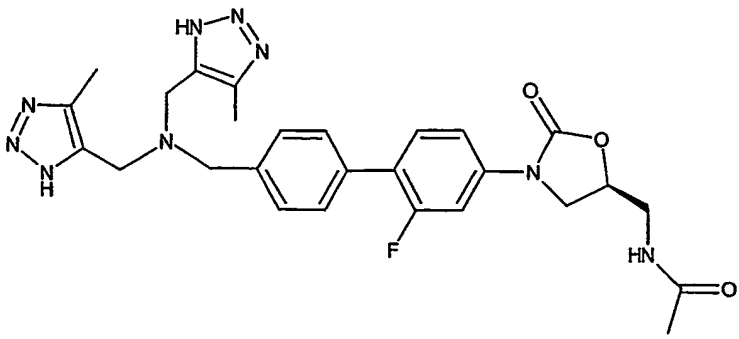
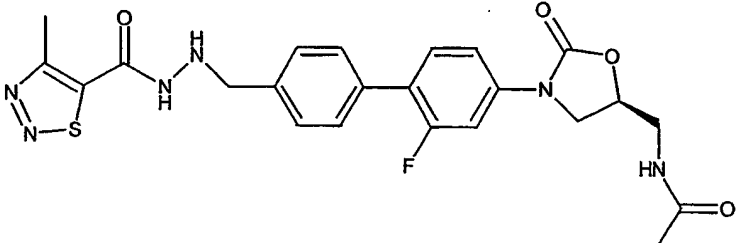
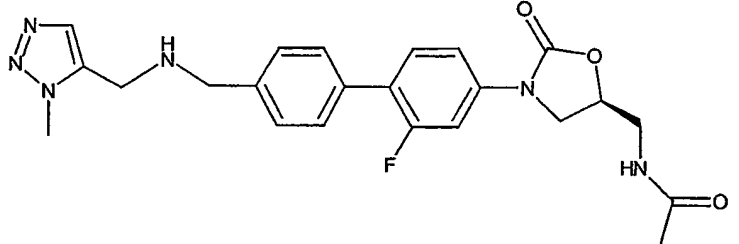
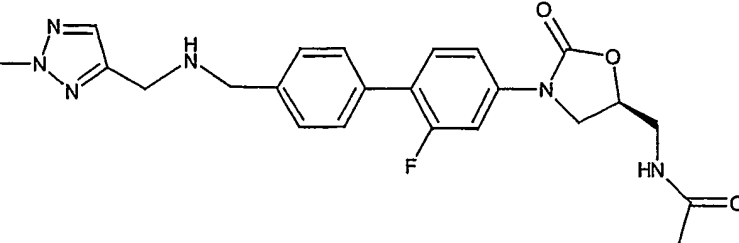
4249	
	N-[3-(4'-{2-[Benzyl-(3-chloro-3,3-difluoro-propyl)-amino]-1-(S)-hydroxy-ethyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4250	
	N-(2-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-yl}-2-(S)-hydroxy-ethyl)-N-benzyl-acetamide
4251	
	N-(3-{4'-[2-(Benzyl-methyl-amino)-1-(S)-hydroxy-ethyl]-2-fluoro-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4252	
	N-{3-[3-Fluoro-4-(6-{[(isoxazol-4-ylmethyl)-amino]-methyl}-pyridin-3-yl)-phenyl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide

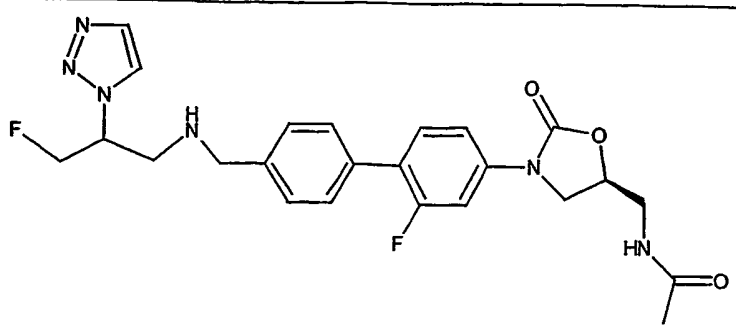
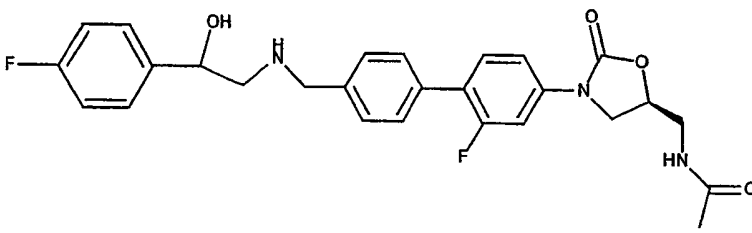
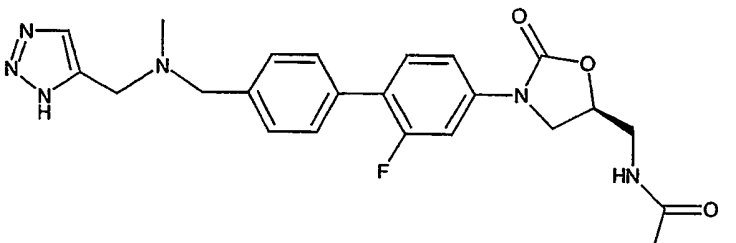
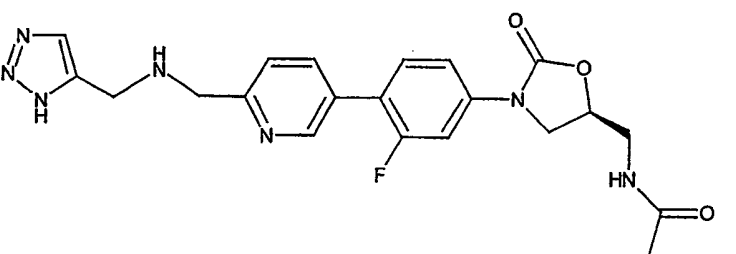
4253	
	N-[3-(4'-{[(3-Bromo-isoxazol-5-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4254	
	N-[3-(2-Fluoro-4'-{2-[(isoxazol-4-ylmethyl)-amino]-1-methoxyimino-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4255	
	N-[3-(2-Fluoro-4'-{1-methoxyimino-2-[(oxazol-4-ylmethyl)-amino]-ethyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4256	
	N-[3-(4'-{[3-(1-Benzyl-1H-[1,2,3]triazol-4-yl)-propylamino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

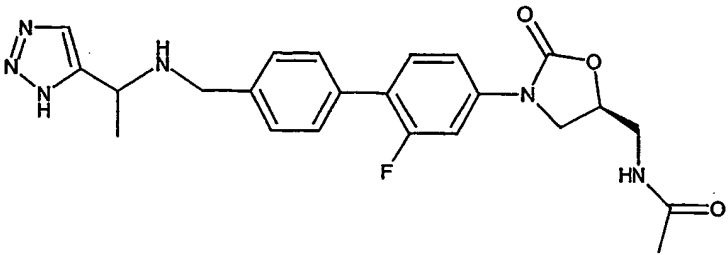
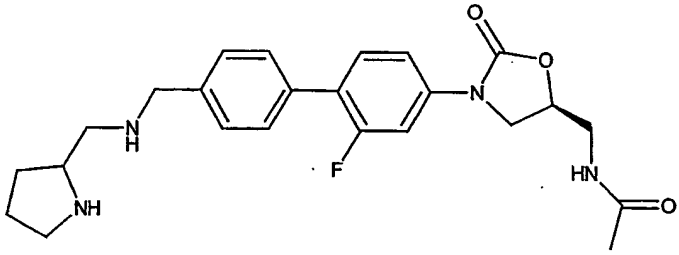
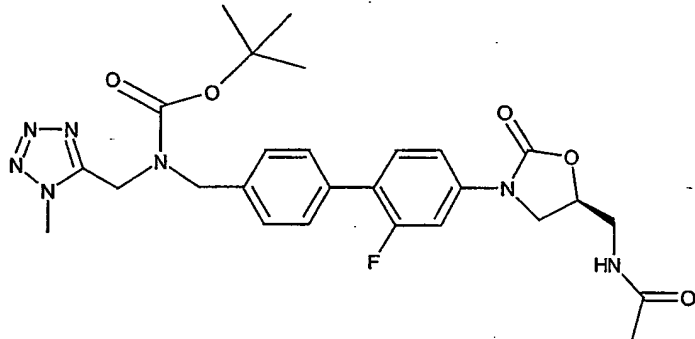
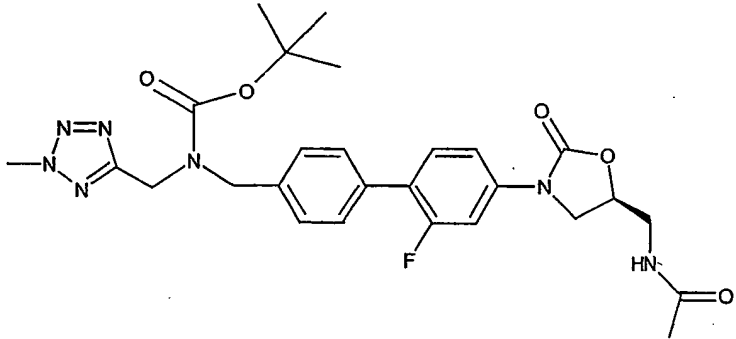
4257	
	N-[3-(2-Fluoro-4'-{[(2-fluoro-pyridin-3-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4258	
	N-[3-(2-Fluoro-4'-{[3-(3H-[1,2,3]triazol-4-yl)-propylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4259	
	N-(3-{2-Fluoro-4'-[(2-pyrrolidin-1-yl-ethylamino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4260	
	N-[3-(3-Fluoro-4-morpholin-4-yl-phenyl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-3-(5-pyrimidin-2-yl-pyridin-2-yl)-propionamide
4261	

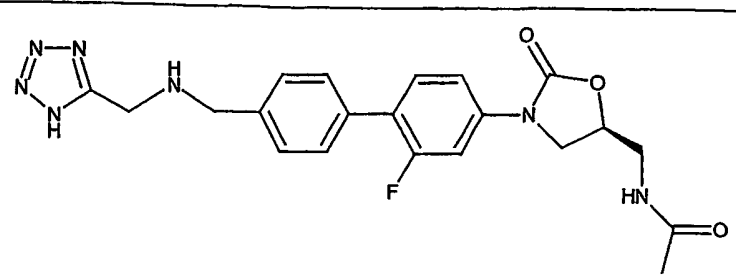
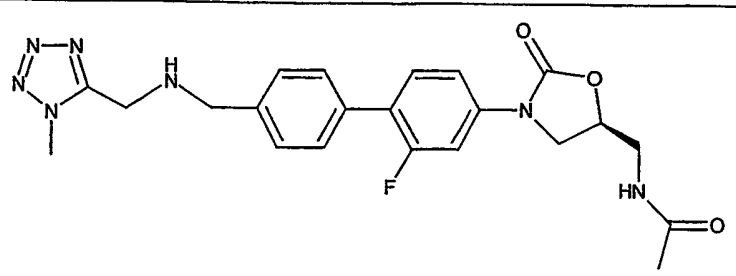
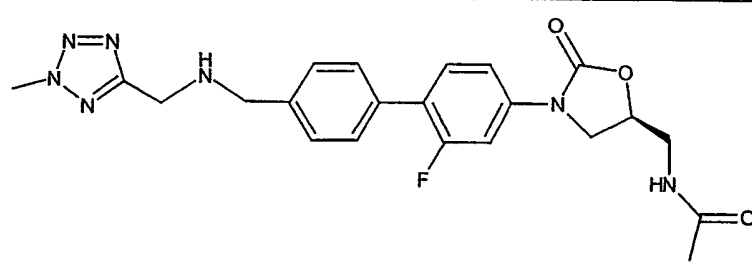
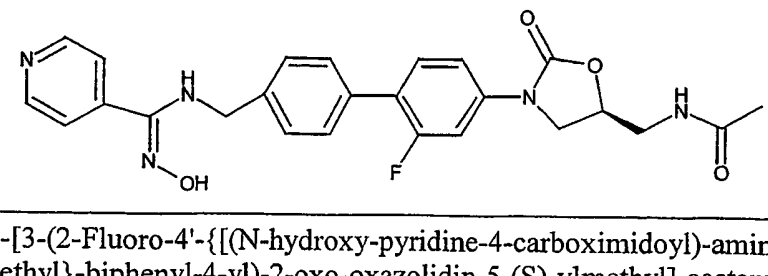
	N-[3-(2-Fluoro-2'-methoxy-4'-{[(pyridin-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4262	
	N-(3-{2-Fluoro-4'-[(2-[1,2,3]triazol-1-yl-ethylamino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4263	
	N-{3-[4'-(Benzyloxyamino-methyl)-2-fluoro-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4264	
	N-(3-{2-Fluoro-4'-[(3-[1,2,3]triazol-1-yl-propylamino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4265	
	N-[3-(4'-{[Benzyloxy-(3-fluoro-propyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

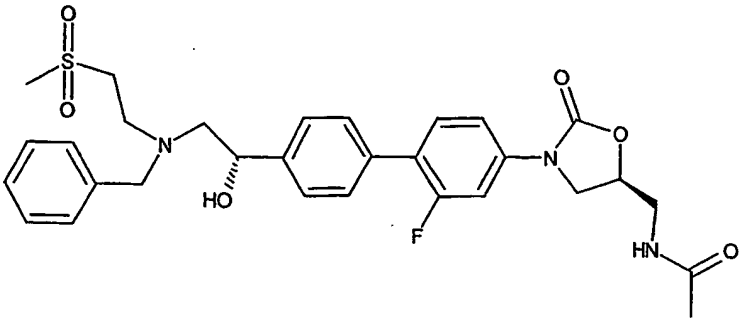
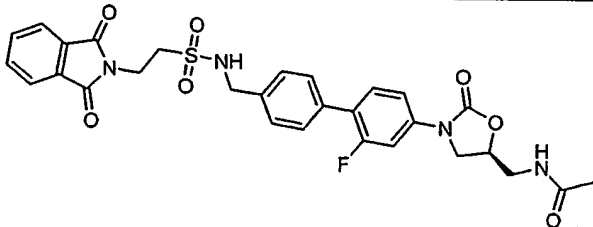
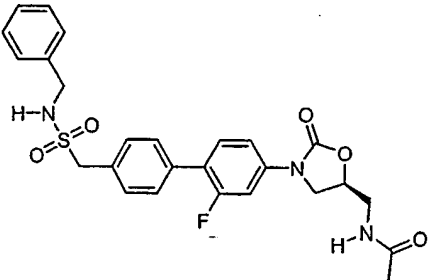
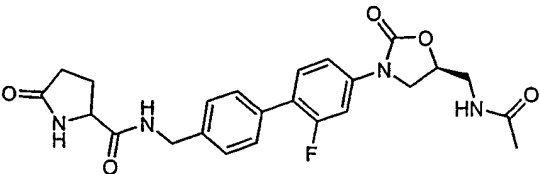
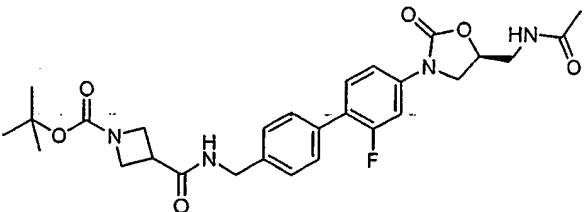
4266	
	N-[3-(2-Fluoro-4'-{[2-(3H-[1,2,3]triazol-4-yl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4267	
	N-[3-(2-Fluoro-4'-{[(3H-[1,2,3]triazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4268	
	3-(2-Fluoro-4'-{[(3H-[1,2,3]triazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-5-(R)-[1,2,3]triazol-1-ylmethyl-oxazolidin-2-one
4269	
	N-[3-(2-Fluoro-4'-{[(5-methyl-3H-[1,2,3]triazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

4270	
	<p>N-[3-(4'-{[Bis-(5-methyl-3H-[1,2,3]triazol-4-ylmethyl)-amino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide</p>
4271	
	<p>N-(3-{2-Fluoro-4'-[N'-(4-methyl-[1,2,3]thiadiazole-carbonyl)hydrazinomethyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide</p>
4272	
	<p>N-[3-(2-Fluoro-4'-{[(3-methyl-3H-[1,2,3]triazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide</p>
4273	
	<p>N-[3-(2-Fluoro-4'-{[(2-methyl-2H-[1,2,3]triazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide</p>

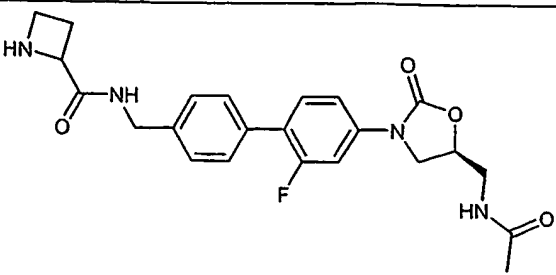
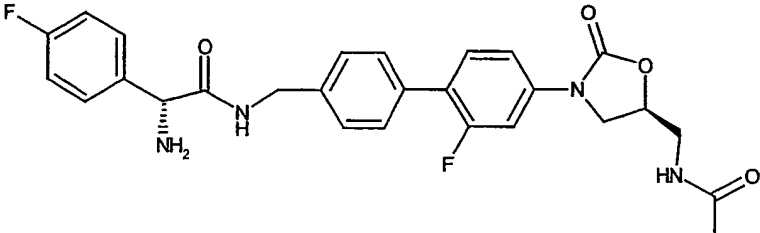
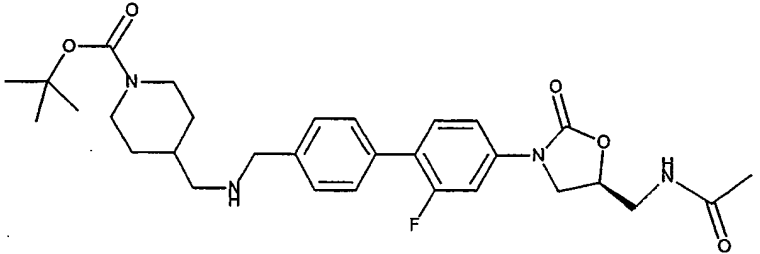
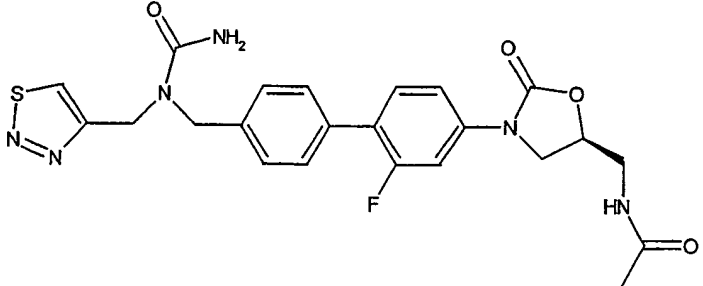
4274	
	N-(3-(2-Fluoro-4'-[(3-fluoro-2-[1,2,3]triazol-1-yl-propylamino)-methyl]-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4275	
	N-[3-(2-Fluoro-4'-{[2-(4-fluoro-phenyl)-2-(R/S)-hydroxy-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4276	
	N-[3-(2-Fluoro-4'-{[methyl-(3H-[1,2,3]triazol-4-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4277	
	N-{3-[3-Fluoro-4-(6-{[(3H-[1,2,3]triazol-4-ylmethyl)-amino]-methyl}-pyridin-3-yl)-phenyl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide

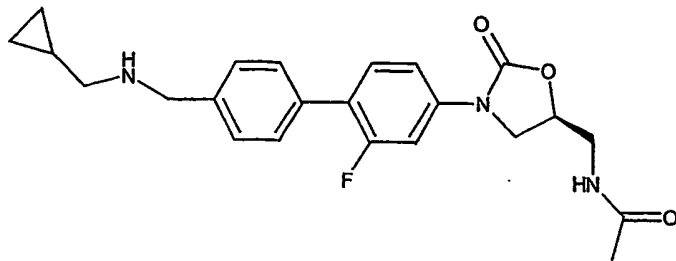
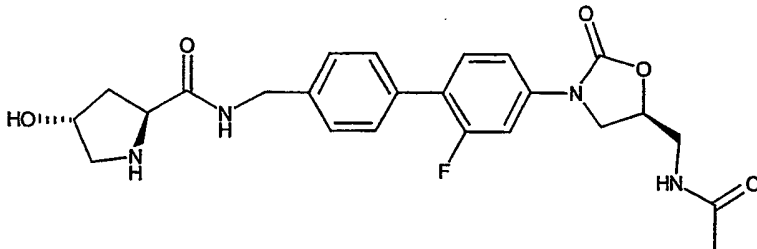
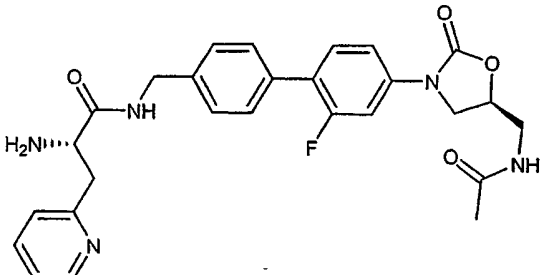
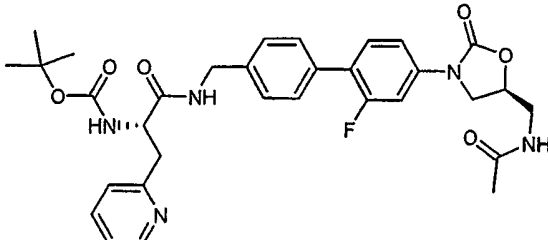
4278	
	<p>N-[3-(2-Fluoro-4'-{[1-(R/S)-(3H-[1,2,3]triazol-4-yl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide</p>
4279	
	<p>N-[3-(2-Fluoro-4'-{[(pyrrolidin-2-(R/S)-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide</p>
4280	
	<p>{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro biphenyl-4-ylmethyl}-(1-methyl-1H-tetrazol-5-ylmethyl)-carbamic acid tert-butyl ester</p>
4281	

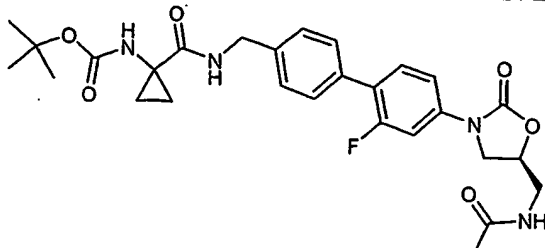
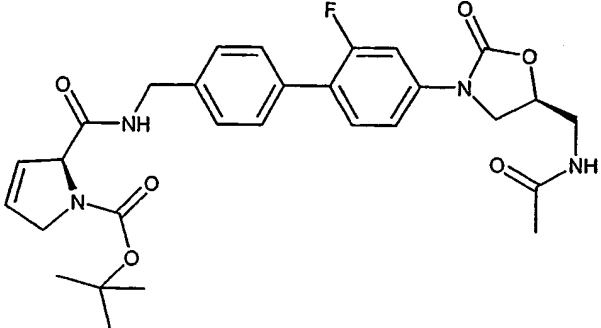
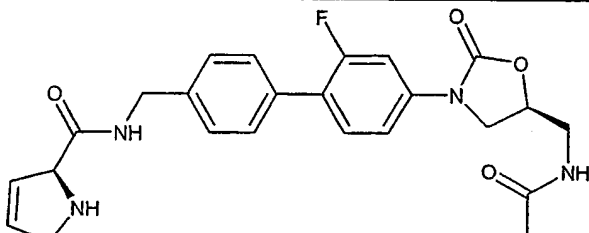
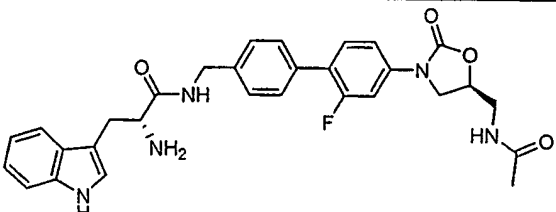
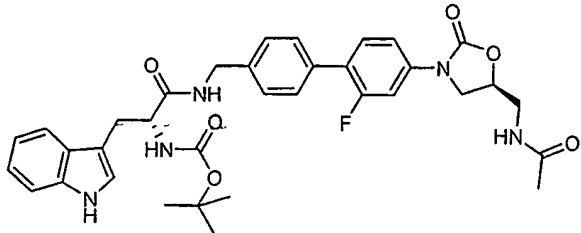
	{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-(2-methyl-2H-tetrazol-5-ylmethyl)-carbamic acid tert-butyl ester
4282	
	N-[3-(2-Fluoro-4'-{[(1H-tetrazol-5-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4283	
	N-[3-(2-Fluoro-4'-{[(1-methyl-1H-tetrazol-5-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4284	
	N-[3-(2-Fluoro-4'-{[(2-methyl-2H-tetrazol-5-ylmethyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4285	
	N-[3-(2-Fluoro-4'-{[(N-hydroxy-pyridine-4-carboximidoyl)-amino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

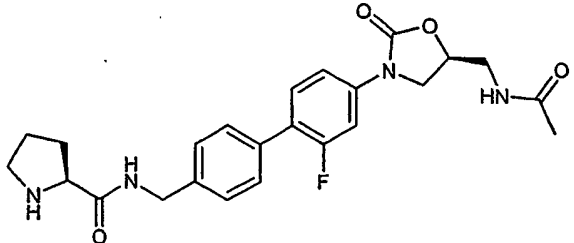
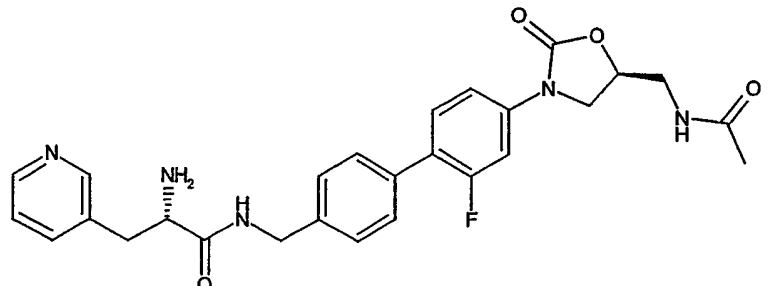
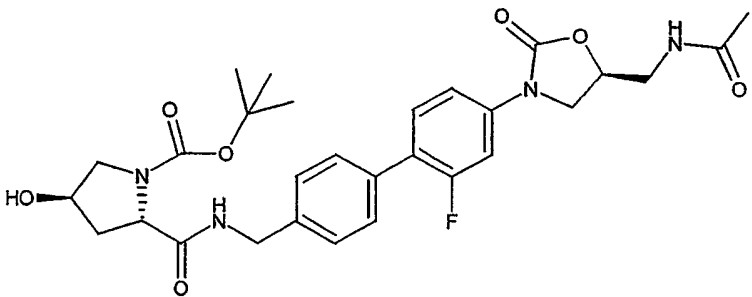
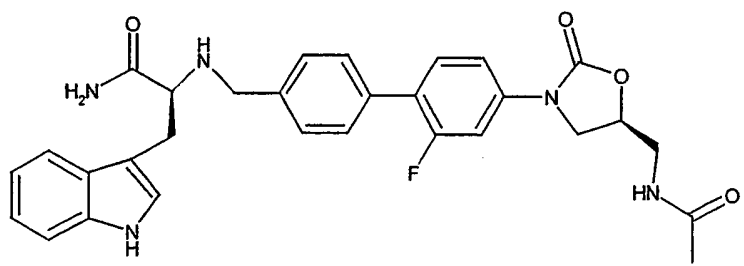
4286	
	N-[3-(4'-{2-[Benzyl-(2-methanesulfonyl-ethyl)-amino]-1-(S)-hydroxy-ethyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4287	
	N-[3-(4'-{2-[(1,3-Dioxo-1,3-dihydro-isoindol-2-yl)-ethanesulfonylamino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
4288	
	N-{3-[4'-(Benzylsulfamoyl-methyl)-2-fluoro-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
4289	
	5-Oxo-pyrrolidine-2-carboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide
4290	

	3-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-carbamoyl)-azetidine-1-carboxylic acid tert-butyl ester
4291	
	Azetidine-3-carboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide
4292	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-2-(R)-amino-3-(3H-imidazol-4-yl)-propionamide
4293	
	2-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amino)-2-pyridin-3-yl-acetamide
4294	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-2-amino-2-pyridin-3-yl-acetamide
4295	

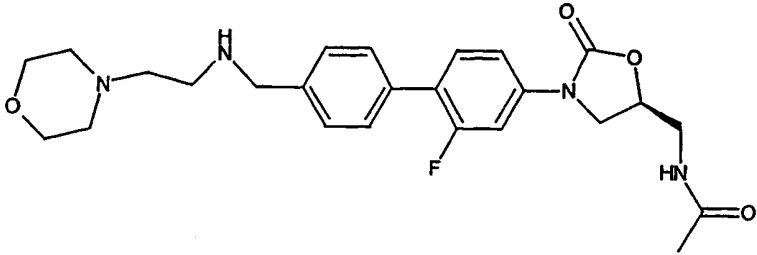
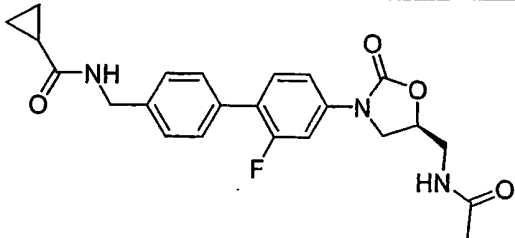
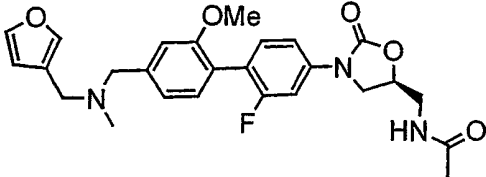
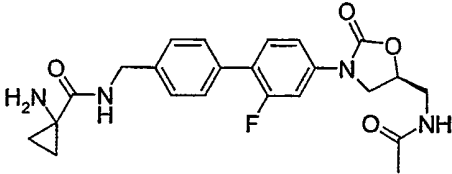
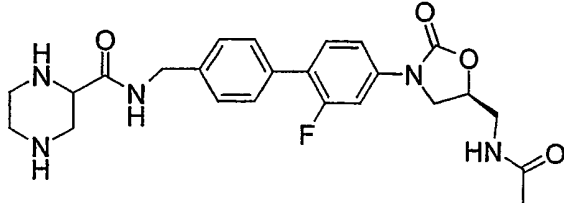
	2-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-carbamoyl)-azetidine-1-carboxylic acid tert-butyl ester
4296	
	Azetidine-2-carboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide
4297	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-2-(R)-amino-2-(4-fluoro-phenyl)-acetamide
4298	
	4-[(4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl)-amino)-methyl]-piperidine-1-carboxylic acid tert-butylester
4299	
	N-{3-[2-Fluoro-4'-(1-[1,2,3]thiadiazol-4-ylmethyl)-ureidomethyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide

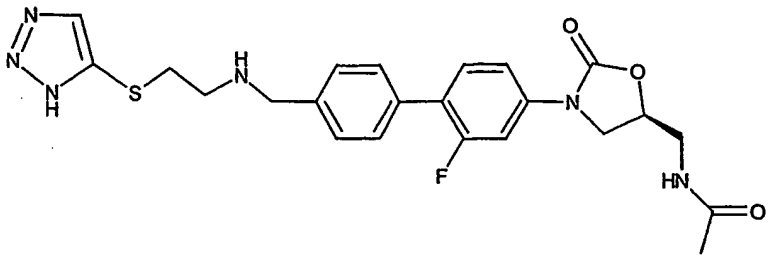
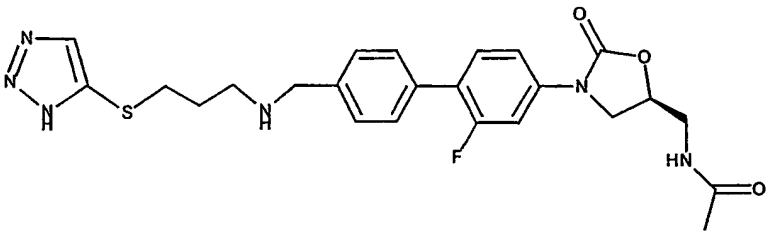
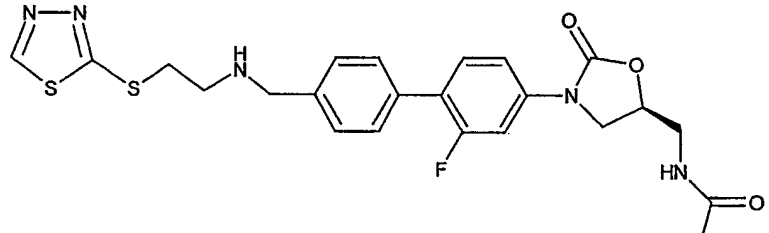
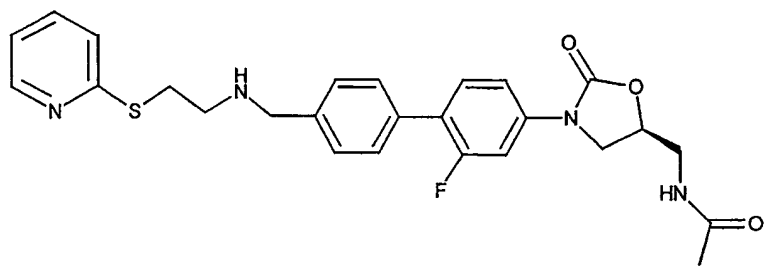
4300	
	N-(3-{4'-[(Cyclopropylmethyl-amino)-methyl]-2-fluoro-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4301	
	4-(R)-Hydroxy-pyrrolidine-2-(S)-carboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide
4302	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-2-(S)-amino-3-pyridin-2-yl-propionamide
4303	
	[1-(S)-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-carbamoyl)-2-pyridin-2-yl-ethyl]-carbamic acid tert-butyl ester

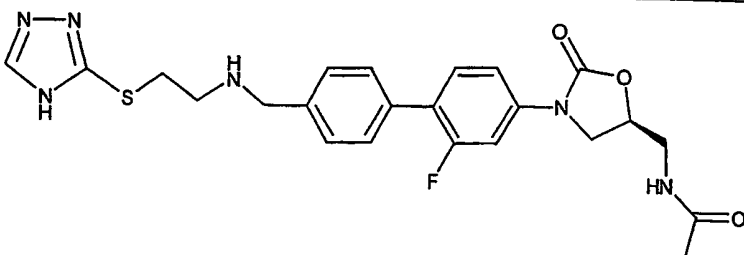
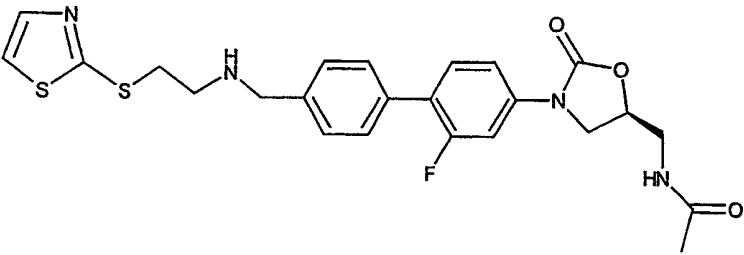
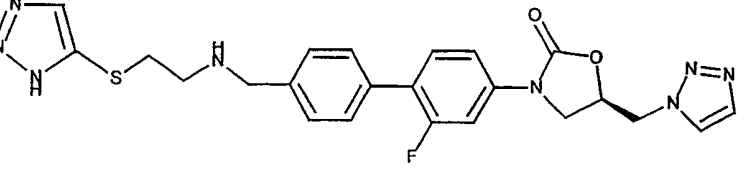
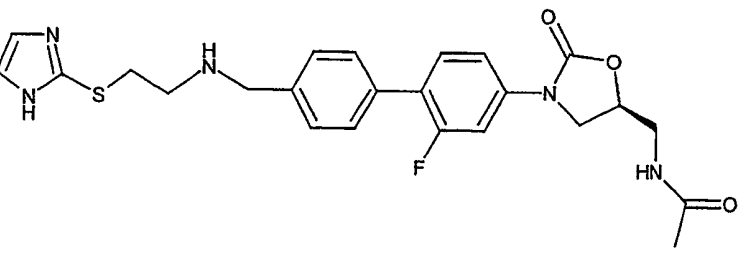
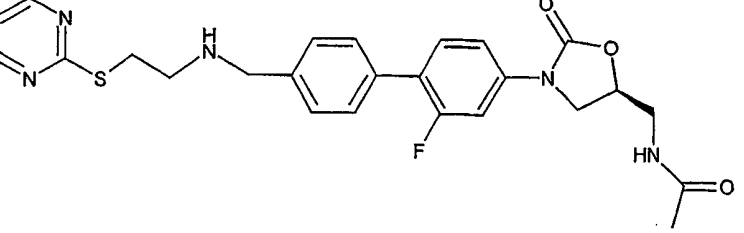
4304	
	[1-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-carbamoyl)-cyclopropyl]-carbamic acid tert-butyl ester
4305	
	2-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-carbamoyl)-2,5-dihydro-pyrrole-1-(S)-carboxylic acid tert-butylester
4306	
	2,5-Dihydro-1H-pyrrole-2-(S)-carboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide
4307	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-2-(R)-amino-3-(1H-indol-3-yl)-propionamide
4308	

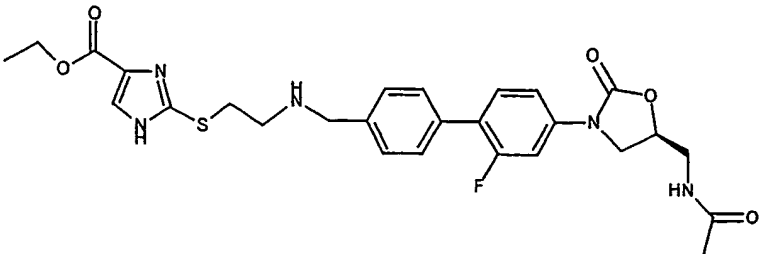
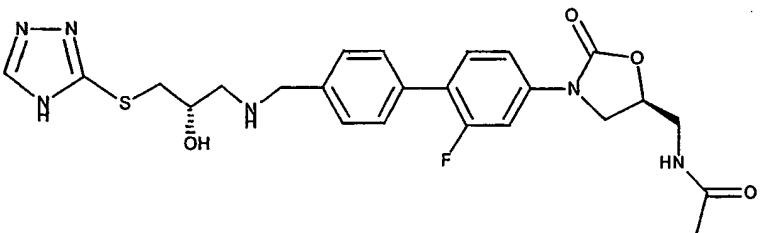
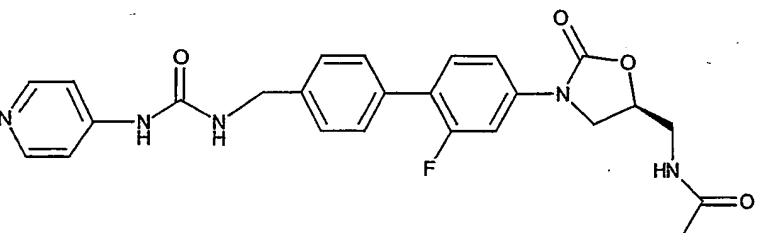
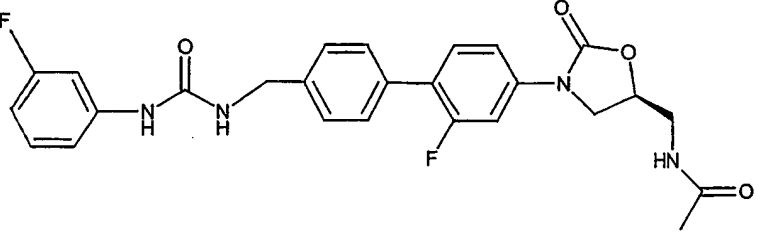
	[1-(R)-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-carbamoyl)-2-(1H-indol-3-yl)-ethyl]-carbamic acid tert-butyl ester
4309	
	Pyrrolidine-2-(S)-carboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide
4310	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-2-(R)-amino-3-pyridin-3-yl-propionamide
4311	
	2-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-carbamoyl)-4-(R)-hydroxy-pyrrolidine-1-(S)-carboxylic acid tert-butyl ester
4312	
	2-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amino)-3-(S)-(1H-indol-3-yl)-propionamide

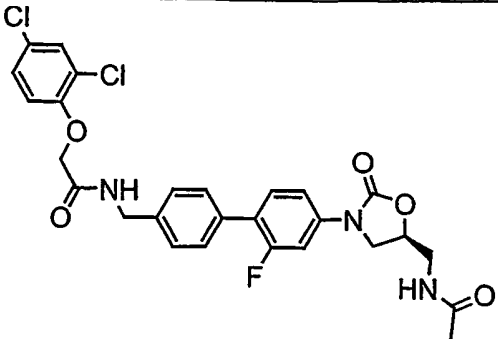
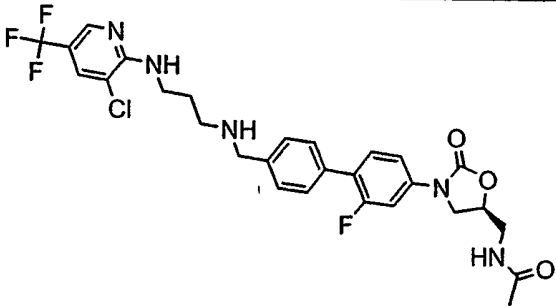
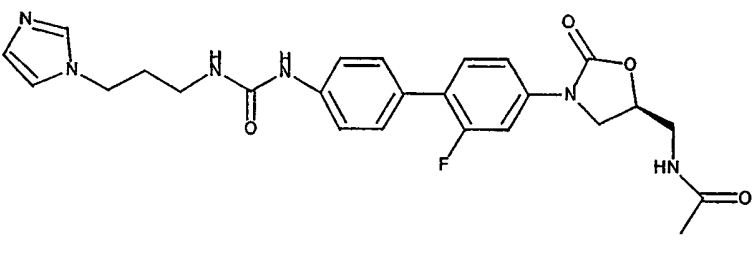
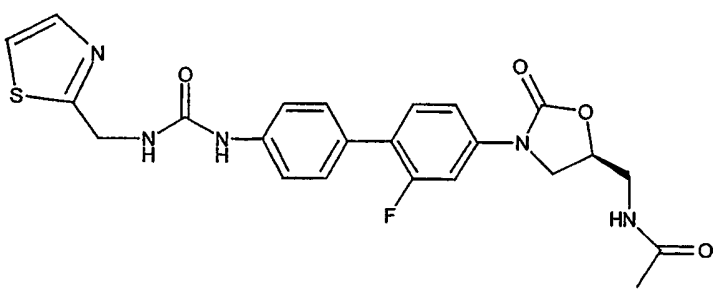
4313	
	2-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amino)-3-(1H-imidazol-4-yl)-propionamide
4314	
	N-(3-{2-Fluoro-4'-[(2-oxo-2-piperazin-1-yl-ethylamino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4315	
	4-[2-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amino)-acetyl]-piperazine-1-carboxylic acid tert-butyl ester
4316	
	N-(3-{2-Fluoro-4'-[(2-morpholin-4-yl-2-oxo-ethylamino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4317	
	3-[(4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl]-amino)-methyl]-pyrrolidine-1-carboxylic acid tert-butyl ester

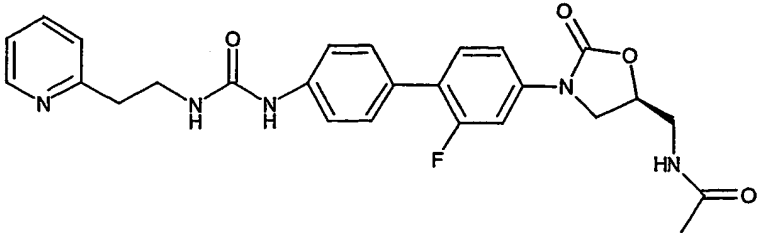
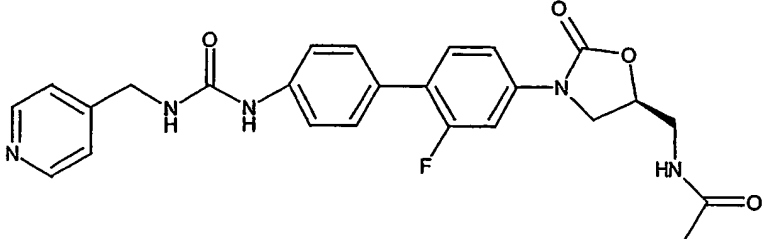
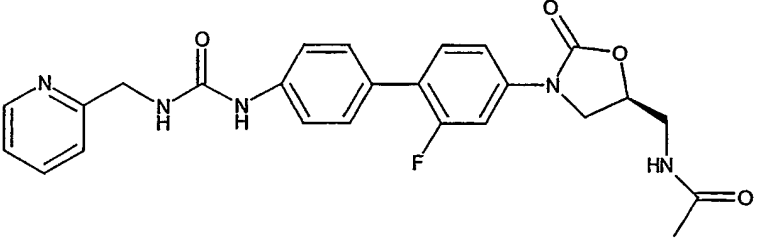
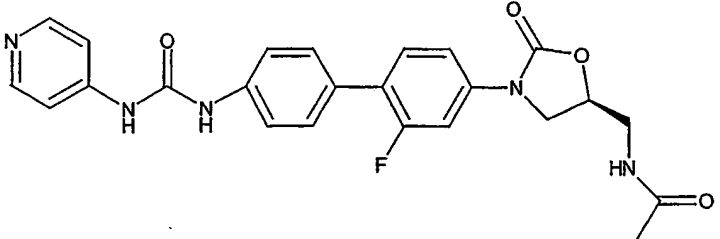
4318	
	N-(3-{2-Fluoro-4'-[(2-morpholin-4-yl-ethylamino)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4319	
	Cyclopropanecarboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide
4320	
	N-(3-{2-Fluoro-4'-[(furan-3-ylmethyl-methyl-amino)-methyl]-2'-methoxy-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
4321	
	1-Amino-cyclopropanecarboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide
4322	
	Piperazine-2-(R/S)-carboxylic acid {4'-[5-(S)-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amide

5001	
	N-[3-(2-Fluoro-4'-{[2-(3H-[1,2,3]triazol-4-ylsulfanyl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
5002	
	N-[3-(2-Fluoro-4'-{[3-(3H-[1,2,3]triazol-4-ylsulfanyl)-propylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
5003	
	N-[3-(2-Fluoro-4'-{[2-([1,3,4]thiadiazol-2-ylsulfanyl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
5004	
	N-[3-(2-Fluoro-4'-{[2-(pyridin-2-ylsulfanyl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide

5005	
	N-[3-(2-Fluoro-4'-{[2-(4H-[1,2,4]triazol-3-ylsulfanyl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
5006	
	N-[3-(2-Fluoro-4'-{[2-(thiazol-2-ylsulfanyl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
5007	
	3-(2-Fluoro-4'-{[2-(3H-[1,2,3]triazol-4-ylsulfanyl)-ethylamino]-methyl}-biphenyl-4-yl)-5-(R)-[1,2,3]triazol-1-ylmethyl-oxazolidin-2-one
5008	
	N-[3-(2-Fluoro-4'-{[2-(1H-imidazol-2-ylsulfanyl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
5009	

	N-[3-(2-Fluoro-4'-{[2-(pyrimidin-2-ylsulfanyl)-ethylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
5010	
	2-[2-({4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-amino)-ethylsulfanyl]-1H-imidazole-4-carboxylic acid ethyl ester
5011	
	N-[3-(2-Fluoro-4'-{[2-(S)-(hydroxy-3-(4H-[1,2,4]triazol-3-ylsulfanyl)-propylamino]-methyl}-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
5012	
	N-(3-{2-Fluoro-4'-[(3-pyridin-4-yl-ureido)-methyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
5013	
	N-(3-{2-Fluoro-4'-[3-(3-fluoro-phenyl)-ureidomethyl]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide

5014	
	N-{4'-[5-(S)-(Acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl}-2-(2,4-dichloro-phenoxy)-acetamide
5015	
	N-[3-(4'-{[3-(3-Chloro-5-trifluoromethyl-pyridin-2-ylamino)-propylamino]-methyl}-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-(S)-ylmethyl]-acetamide
6001	
	N-(3-{2-Fluoro-4'-[3-(3-imidazol-1-yl-propyl)-ureido]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
6002	
	N-{3-[2-Fluoro-4'-(3-thiazol-2-ylmethyl-ureido)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide

6003	
	N-(3-{2-Fluoro-4'-[3-(2-pyridin-2-yl-ethyl)-ureido]-biphenyl-4-yl}-2-oxo-oxazolidin-5-(S)-ylmethyl)-acetamide
6004	
	N-{3-[2-Fluoro-4'-(3-pyridin-4-ylmethyl-ureido)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
6005	
	N-{3-[2-Fluoro-4'-(3-pyridin-2-ylmethyl-ureido)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide
6006	
	N-{3-[2-Fluoro-4'-(3-pyridin-4-yl-ureido)-biphenyl-4-yl]-2-oxo-oxazolidin-5-(S)-ylmethyl}-acetamide

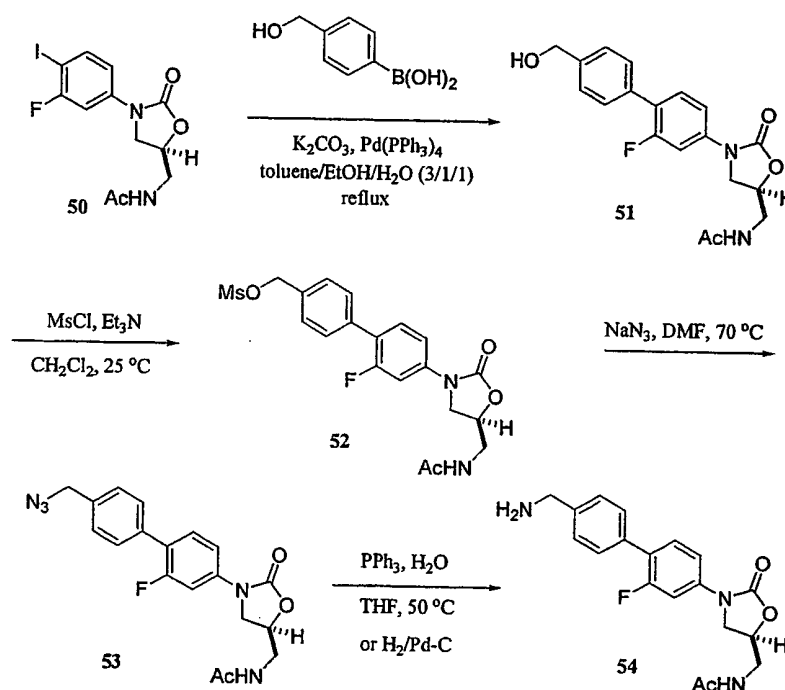
Nuclear magnetic resonance (NMR) spectra were obtained on a Bruker Avance 300 or Avance 500 spectrometer, or in some cases a GE-Nicolet 300 spectrometer. Common reaction solvents were either high performance liquid chromatography (HPLC) grade or American Chemical Society (ACS) grade, and anhydrous as obtained from the manufacturer unless

otherwise noted. "Chromatography" or "purified by silica gel" refers to flash column chromatography using silica gel (EM Merck, Silica Gel 60, 230-400 mesh) unless otherwise noted.

Example 1 – Synthesis of Biaryl Precursors

5 Scheme 1 depicts the synthesis of various biaryl intermediates useful in producing compounds of the present invention. Known iodoaryl oxazolidinone intermediate **50** (see U.S. Patent Nos. 5,523,403 and 5,565,571) is coupled to a substituted aryl boronic acid (the Suzuki reaction) to produce biaryl alcohol **51**. Mesylate **52**, azide **53**, and amine **54** are then synthesized using chemistry well known to those skilled in the art.

10 Scheme 1



Synthesis of alcohol **51**

A suspension of *N*-[3-(3-fluoro-4-iodo-phenyl)-2-oxo-oxazolidin-5-ylmethyl]-acetamide **50** (14.0 g, 37 mmol) in toluene (120 mL) was treated with 4-(hydroxymethyl)phenylboronic acid (7.87 g, 51.8 mmol, 1.4 equiv), potassium carbonate (K_2CO_3 , 15.32 g, 111 mmol, 3.0 equiv), ethanol (EtOH, 40 mL), and H₂O (40 mL) at 25 °C, and the resulting mixture was degassed three times under a steady stream of argon at 25 °C.

Tetrakis(triphenylphosphine)palladium ($Pd(PPh_3)_4$, 2.14 g, 1.85 mmol, 0.05 equiv) was subsequently added to the reaction mixture, and the resulting reaction mixture was degassed

three times again before being warmed to gentle reflux for 6 h. When thin layer chromatography (TLC) and HPLC showed the coupling reaction was complete, the reaction mixture was cooled to room temperature before being treated with H₂O (240 mL). The resulting mixture was then stirred at room temperature for 10 min before being cooled to 0-5 °C for 1 h. The solid precipitates were collected by filtration, washed with H₂O (2 x 100 mL) and 20% ethyl acetate (EtOAc)/hexane (2 X 50 mL), and dried *in vacuo*. The crude desired *N*-[3-(2-Fluoro-4'-hydroxymethyl-biphenyl-4-yl)-2-oxo-oxazolidin-5-ylmethyl]-acetamide **51** (12.50 g, 94% yield) was obtained as off-white solids. This material was found to be essentially pure by HPLC and ¹H NMR and was directly used in the subsequent reaction without further purification. ¹H NMR (300 MHz, DMSO-*d*₆) δ 1.76 (s, 3H, COCH₃), 3.35 (t, 2H, *J* = 5.4 Hz), 3.69 (dd, 1H, *J* = 6.4, 9.2 Hz), 4.08 (t, 1H, *J* = 9.1 Hz), 4.46 (d, 2H, *J* = 5.7 Hz, CH₂OH), 4.68 (m, 1H), 5.16 (t, 1H, *J* = 5.7 Hz, OH), 7.25 – 7.52 (m, 7H, aromatic-*H*), 8.18 (t, 1H, *J* = 5.8 Hz, NHCOCH₃). LCMS (ESI) *m/e* 359 (M + H)⁺.

Synthesis of mesylate **52**

A suspension of **51** (12.49 g, 34.90 mmol) in methylene chloride (CH₂Cl₂, 150 mL) was treated with triethylamine (Et₃N, 7.07 g, 9.7 mL, 70 mmol, 2.0 equiv) at 25 °C, and the resulting mixture was cooled to 0-5 °C before being treated dropwise with methanesulfonyl chloride (4.80 g, 3.24 mL, 41.9 mmol, 1.2 equiv) at 0-5 °C. The resulting reaction mixture was subsequently stirred at 0-5 °C for 2 h. When TLC and HPLC showed the reaction was complete, the reaction mixture was treated with H₂O (100 mL) at 0-5 °C. The mixture was then concentrated *in vacuo* to remove most of the CH₂Cl₂, and the resulting slurry was treated with H₂O (150 mL). The mixture was stirred at room temperature for 10 min before being cooled to 0-5 °C for 30 min. The solid precipitates were collected by filtration, washed with H₂O (2 x 100 mL) and 20% EtOAc/hexane (2 X 50 mL), and dried *in vacuo*. The crude desired methanesulfonic acid 4'-[5-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-ylmethyl ester **52** (11.84 g, 78% yield) was obtained as off-white solids, which by TLC and HPLC was found to be essentially pure and was directly used in the subsequent reaction without further purification. LCMS (ESI) *m/e* 437 (M + H)⁺.

Synthesis of azide **53**

A solution of **52** (9.27 g, 21.26 mmol) in anhydrous *N,N*-dimethylformamide (DMF, 50 mL) was treated with sodium azide (NaN₃, 5.53 g, 85.04 mmol, 4.0 equiv) at 25 °C, and the resulting reaction mixture was warmed to 70-80 °C for 4 h. When TLC and HPLC showed the

reaction was complete, the reaction mixture was cooled to room temperature before being treated with H₂O (150 mL). The resulting mixture was stirred at room temperature for 10 min before being cooled to 0–5 °C for 1 h. The solid precipitates were collected by filtration, washed with H₂O (2 x 100 mL) and 20% EtOAc/hexane (2 X 50 mL), and dried *in vacuo*. The crude desired *N*-[3-(4'-azidomethyl-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-ylmethyl]-acetamide **53** (7.16 g, 88% yield) was obtained as off-white solids. The material was found to be essentially pure by TLC and HPLC and was directly used in the subsequent reaction without further purification. LCMS (ESI) *m/e* 384 (M + H)⁺.

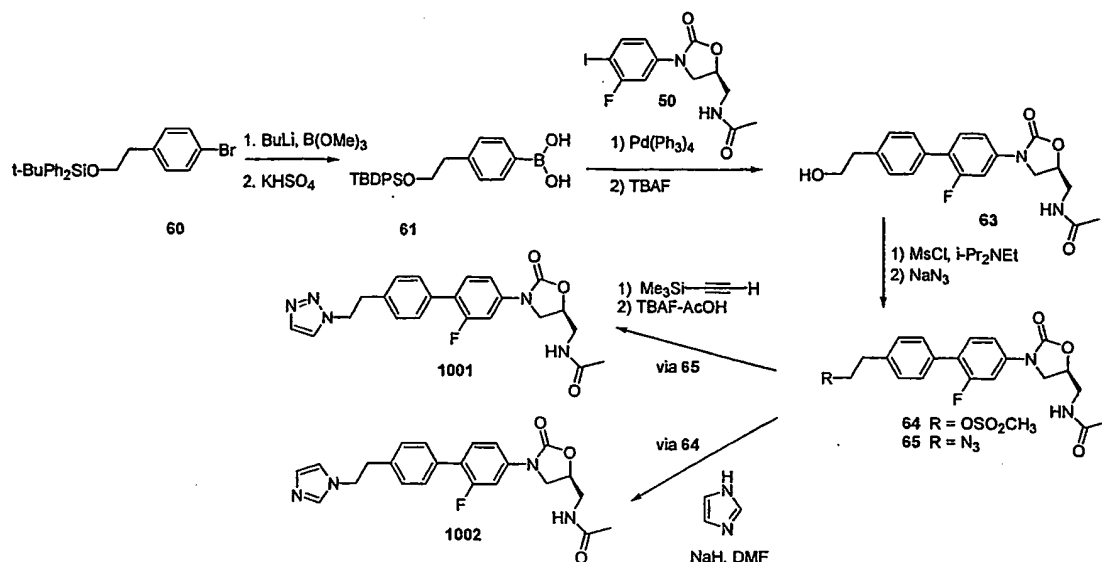
Synthesis of amine **54**

A solution of **53** (7.16 g, 18.69 mmol) in tetrahydrofuran (THF) (100 mL) was treated with triphenylphosphine (PPh₃, 5.88 g, 22.43 mmol, 1.2 equiv) and H₂O (3.6 g, 3.6 mL, 0.2 mmol, 11.0 equiv) at 25 °C, and the resulting reaction mixture was warmed to 50–55 °C for 12 h. When TLC and HPLC showed the reduction reaction was complete, the reaction mixture was cooled to room temperature before the solvents were removed *in vacuo*. The residue was directly purified by flash column chromatography (0–15% MeOH-CH₂Cl₂ gradient elution) to afford the desired *N*-[3-(4'-Aminomethyl-2-fluoro-biphenyl-4-yl)-2-oxo-oxazolidin-5-ylmethyl]-acetamide **54** (5.82 g, 87% yield) as off-white crystals, which were of sufficient purity to be directly used in subsequent reactions. ¹H NMR (300 MHz, DMSO-*d*₆) δ 1.85 (s, 3H, COCH₃), 3.04 (br. s, 2H, NH₂), 3.44 (t, 2H, *J* = 5.4 Hz), 3.78 (m, 3H), 4.18 (t, 1H, *J* = 9.1 Hz), 4.77 (m, 1H), 7.25 – 7.60 (m, 7H, aromatic-*H*), 8.20 (t, 1H, *J* = 5.8 Hz, NHCOCH₃). LCMS (ESI) *m/e* 359 (M + 2H)²⁺.

Example 2 - Synthesis of Triazole **1001** and Imidazole **1002**

Scheme 2 illustrates the synthesis of triazole **1001** and imidazole **1002**. Aryl bromide **60** was converted to boronic acid **61** which was used in a Suzuki coupling with aryl iodide **50** to afford alcohol **63** after desilylation. The alcohol was converted to mesylate **64** and then to azide **65**. The cycloaddition of azide **65** with trimethylsilylacetylene followed by desilylation afforded triazole **1001**. Alkylation of mesylate **64** with imidazole yielded compound **1002**.

Scheme 2



Synthesis of bromide 60

To a solution of 4-bromophenethyl alcohol (5.60 g, 27.9 mmol), imidazole (3.80 g, 55.7 mmol) and a catalytic amount of 4-dimethylaminopyridine (DMAP) in DMF (55 mL) was added *t*-butyldiphenylchlorosilane (TBDPSCl, 7.20 mL, 27.9 mmol) at 0 °C and the mixture was stirred at ambient temperature for 72 h. The reaction was quenched with ice cold water (50 mL) and extracted with ether (4 x 50 mL). The combined ethereal layer was washed with water (4 x 100 mL), dried over anhydrous sodium sulfate (Na₂SO₄), concentrated and purified by flash chromatography (2% ethyl acetate in hexanes) to yield 10.6 g of 60.

Synthesis of boronic acid 61

To a solution of 60 (10.5 g, 24.0 mmol) in THF (50 mL) was added *n*-butyl lithium (*n*-BuLi, 2.5M in hexane, 11.5 mL, 28.8 mmol) at -78 °C and the mixture was stirred for 1 h before the addition of trimethyl borate (3.54 mL, 31.2 mmol). The solution was then stirred overnight at ambient temperature and quenched with 1M potassium hydrogen sulfate (KHSO₄, 25 mL). The resulting mixture was extracted with CH₂Cl₂ (3 x 50 mL), washed with brine (3 x 100 mL), dried (anhydrous Na₂SO₄), concentrated and purified by flash chromatography (25% ethyl acetate in hexanes) to yield 5 g of boronic acid 61 as mixture of acid and cyclic anhydrides.

Synthesis of alcohol 63

To a mixture of boronic acid 61 (4.7 g, 11.7 mmol), known oxazolidinone 50 (4.00 g, 10.6 mmol; *see* U.S. Patent Nos. 5,523,403 and 5,565,571), potassium carbonate (K₂CO₃, 4.40

g, 31.8 mmol) and $\text{Pd}(\text{PPh}_3)_4$ (0.613 g, 5 mol%) was added toluene (90 mL), ethanol (30 mL) and H_2O (30 mL). The reaction mixture was refluxed overnight under argon atmosphere, concentrated and redissolved in CH_2Cl_2 (100 mL). The organic phase was washed with brine solution (2 x 100 mL), dried (anhydrous Na_2SO_4), concentrated and used for the next step
5 without further purification. To a solution of this crude material in THF (70 mL) was added tetrabutylammonium fluoride (TBAF, 20 mL, 20 mmol) and the mixture was stirred overnight at ambient temperature. The reaction mixture was concentrated and washed with water (4 x 100 mL) to yield 3.5 g of **63**. LCMS (ESI) m/z 373 (M+H).

Synthesis of mesylate **64** and azide **65**

10 To a solution of **63** (1.0 g, 2.7 mmol) in CH_2Cl_2 (15 mL), DMF (4 mL) and *N,N*-diisopropylethylamine (Hunig's base, 0.75 mL, 4.05 mmol) was added methanesulfonyl chloride (0.32 mL, 2.7 mmol) at 0 °C. After 2 h the reaction mixture was poured into CH_2Cl_2 (150 mL) and the organic layer was washed with water (3 x 100 mL), dried, concentrated to afford **64** as a solid. The crude solid **64** thus obtained was heated with NaN_3 (0.35 g, 5.4
15 mmol) at 90 °C overnight. The reaction mixture was poured into ethyl acetate (100 mL). The ethyl acetate layer was washed with water (3 x 50 mL), dried and concentrated to yield 1.1 g of pure azide **65**. LCMS (ESI) m/z 398 (M+H).

Synthesis of triazole **1001**

A solution of azide **65** (100 mg, 0.252 mmol) and trimethylsilylacetylene (0.072 mL, 0.504 mmol) in DMF (3 mL) was heated at 90 °C until the azide was consumed. The reaction
20 mixture was concentrated and treated with TBAF (1 mL, 1 mmol) and acetic acid (0.028 mL, 0.504 mmol) in THF (3 mL). The solution was stirred for 72 h and concentrated. The crude product was purified by flash chromatography using 4% methanol (MeOH) in CH_2Cl_2 to yield 85 mg of **1001**. LCMS (ESI) m/z 424 (M+H).

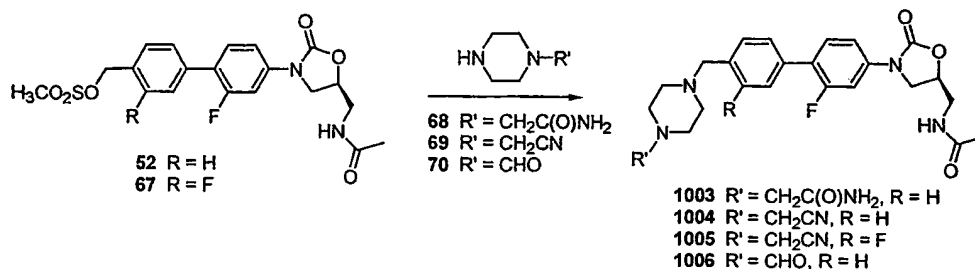
25 Synthesis of imidazole **1002**

To a solution of imidazole (70 mg, 1.0 mmol) in DMF (5 mL) was added sodium hydride (NaH, 60%, 41 mg, 1 mmol) at 0 °C and the mixture was stirred for 30 minutes before the addition of mesylate **64** (114 mg, 0.250 mmol). The resulting solution was heated to 80 °C for 3h, concentrated and purified by flash chromatography (5% MeOH in CH_2Cl_2). After
30 trituration with ether, the residue afforded 40 mg of **1002**. LCMS (ESI) m/z 423 (M+H).

Example 3 - Synthesis of Piperazines 1003-1006

Scheme 3 illustrates the synthesis of compounds **1003-1006**. Mesylate **52** served as alkylating agent for piperazine intermediates **68**, **69** and **70** to afford compounds **1003**, **1004** and **1006** respectively. Mesylate **67** was employed to alkylate piperazine intermediate **69** to provide compound **1005**.

Scheme 3

**Synthesis of mesylate 67**

Mesylate **67** was synthesized by coupling iodide **50** and 4-formyl-3-fluorophenylboronic acid following the procedure described above for the synthesis of *N*-[3-(2-fluoro-4'-hydroxymethyl-biphenyl-4-yl)-2-oxo-oxazolidin-5-ylmethyl]-acetamide (see Example 1). The biaryl aldehyde obtained (1.0 g, 2.67 mmol) was suspended in 40 mL methanol and the mixture was cooled to 0°C. Sodium borohydride (0.112 g, 2.943 mmol) was added, and the mixture was stirred for 50 min. Water was added (20 mL), and after stirring another 20 min the mixture was partitioned between methylene chloride and brine. The aqueous phase was extracted twice with methylene chloride. The aqueous phase was acidified to pH 7, then extracted twice with methylene chloride. The combined organic phase was washed with brine, dried over Na_2SO_4 , and concentrated. The crude material was azeotroped with toluene to afford the expected alcohol (900 mg).

The above alcohol (900 mg) was dissolved in methylene chloride (20 mL), DMF (13 mL) and Hunig's base (1.23 mL) and the mixture was cooled to 0°C. Methanesulfonyl chloride (557 μL , 7.20 mmol) was added and the mixture was stirred for 1.5 h at 0°C. LCMS indicated a mixture of desired mesylate and some of the corresponding benzyl chloride. The mixture was stirred for another 30 min and then concentrated. The residue was treated with 400 mL water, and the precipitate was filtered and washed with water. Drying under vacuum overnight yielded 750 mg crude mesylate **67** (as a mixture with some of the corresponding chloride).

Synthesis of piperazine 68

A solution of *tert*-butyl-1-piperazine carboxylate (1 g, 5.4 mmol), bromoacetamide (820 mg, 5.94 mmol) and Hunig's base (1.2 mL, 7.2 mmol) in a mixture of CH₂Cl₂ (10 mL) and MeOH (10 mL) was heated to reflux for 4 h. The reaction mixture was concentrated and the crude product thus obtained was purified by flash chromatography (19 :1 :0.01 CH₂Cl₂/MeOH/ NH₄OH) to yield 1.3 g of pure BOC-protected piperazinyl acetamide. To a solution of the acetamide (250 mg, 1 mmol) in CH₂Cl₂ (10 mL) was added trifluoroacetic acid (TFA, 5 mL) at 0°C and the mixture was stirred at that temperature for 2 h. The reaction mixture was concentrated to yield **68** which was used for subsequent reactions without further purification.

Synthesis of piperazine 69

A solution of *tert*-butyl-1-piperazine carboxylate (1 g, 5.4 mmol), bromoacetonitrile (0.5 mL, 5.94 mmol) and Hunig's base (1.2 mL, 7.2 mmol) in a mixture of CH₂Cl₂ (10 mL) and MeOH (10 mL) was stirred at ambient temperature for 4 h. The reaction mixture was concentrated and the crude product thus obtained was purified by flash chromatography (19:1:0.01 CH₂Cl₂/MeOH/NH₄OH) to yield 1.3 g of pure BOC-protected piperazinyl acetonitrile. To a solution of the piperazinyl acetonitrile (300 mg, 1.3 mmol) in CH₂Cl₂ (10 mL) was added TFA (5 mL) at 0°C and the mixture was stirred at that temperature for 2 h. The reaction mixture was concentrated to yield **69** which was used for subsequent reactions without further purification.

Synthesis of compound 1003

A solution of mesylate of **52** (138 mg, 0.320 mmol) and **68** (~1 mmol) in Hunig's base (2 mL) and DMF (8 mL) was heated to 90°C for 2 h. Then the solution was concentrated and purified by flash chromatography over silica gel (20:1:0.01 CH₂Cl₂/MeOH/NH₄OH) to yield **1003**. LCMS (ESI) *m/z* 484 (M + H)⁺.

Synthesis of compound 1004

Compound **1004** was synthesized from mesylate **52** and piperazine intermediate **69** in the same manner as described above for the synthesis of compound **1003**. LCMS (ESI) *m/z* 466 (M + H)⁺.

Synthesis of compound 1005

Compound **1005** was synthesized from mesylate **67** and piperazine intermediate **69** in the same manner as described above for the synthesis of compound **1003**. LCMS (ESI) m/z 484 ($M + H$)⁺.

5 Synthesis of compound 1006

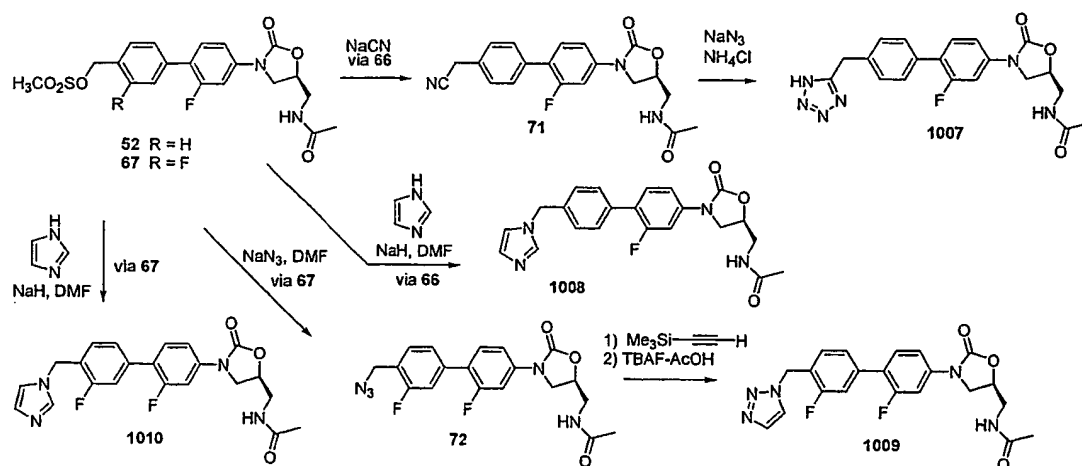
Compound **1006** was synthesized from mesylate **52** and available piperazine intermediate **70** in the same manner as described above for the synthesis of compound **1003**. LCMS (ESI) m/z 455 ($M + H$)⁺.

Example 4 - Synthesis of Compounds 1007-1010

10 Scheme 4 illustrates the synthesis of compounds **1007-1010**. Mesylate **52** was converted to nitrile **71**, which was subsequently transformed to tetrazole **1007**. Mesylate **52** served as alkylating agent for the anion derived from imidazole to afford imidazole derivative **1008**. Mesylate **67** was converted to azide **72**, which was then subsequently converted to triazole **1009**. Mesylate **67** served as alkylating agent for the anion derived from imidazole to afford imidazole derivative **1010**.

15

Scheme 4



Synthesis of tetrazole 1007

To a solution of mesylate **52** (2.0 g, 4.6 mmol) in DMF (30 mL) was added sodium cyanide (NaCN, 0.45 g, 9.2 mmol) and the mixture was heated to 70°C for 3 h. The reaction mixture was cooled to ambient temperature and poured into water (800 mL). The solid thus

20

obtained was filtered and passed through a small bed of silica gel (CH_2Cl_2 : MeOH = 12:1) to yield 1.8 g of nitrile **71**. LCMS (ESI) m/z 368 ($\text{M} + \text{H}$)⁺.

A mixture of **71** (100 mg, 0.272 mmol), NaN_3 (40 mg, 0.598 mmol) and ammonium chloride (NH_4Cl , 32 mg, 0.598 mmol) in DMF (2 mL) was heated to 90°C for 3 days. The reaction mixture was concentrated and purified by flash chromatography (10% MeOH in CH_2Cl_2) to yield 35.6 mg of tetrazole **1007**. LCMS (ESI) m/z 411 ($\text{M} + \text{H}$)⁺.

Synthesis of imidazole **1008**

To a solution of imidazole (37.4 mg, 0.550 mmol) in DMF (5 mL) was added NaH (60%, 20 mg, 0.50 mmol) at 0°C and the mixture was stirred for 30 minutes before the addition of mesylate **52** (200 mg, 0.459 mmol). The resulting solution was heated to 60°C for 2 h and poured into water (75 mL). The aqueous suspension was extracted with 10% MeOH in CH_2Cl_2 (3 x 75 mL) and the combined organic layer was washed with saturated NH_4Cl solution (2 x 100 mL). The organic layer was dried (anhydrous Na_2SO_4), concentrated and triturated with ether to yield 170 mg of imidazole **1008**. LCMS (ESI) m/z 409 ($\text{M} + \text{H}$)⁺.

Synthesis of azide **72**

Crude mesylate **67** (100 mg, 0.224 mmol; as a mixture with some corresponding benzyl chloride) was dissolved in DMF (10 mL) and sodium azide (114.6 mg, 1.762 mmol) was added. The mixture was stirred at room temperature for 14 h, and then partitioned between ethyl acetate and water. The organic phase was washed with water, dried over Na_2SO_4 , and concentrated to provide azide **72** as a solid (190 mg).

Synthesis of triazole **1009**

Compound **1009** was synthesized from azide **72** and trimethylsilylacetylene in the same manner as described above for the synthesis of triazole **1001**. LCMS (ESI) m/z 428 ($\text{M} + \text{H}$)⁺.

Synthesis of imidazole **1010**

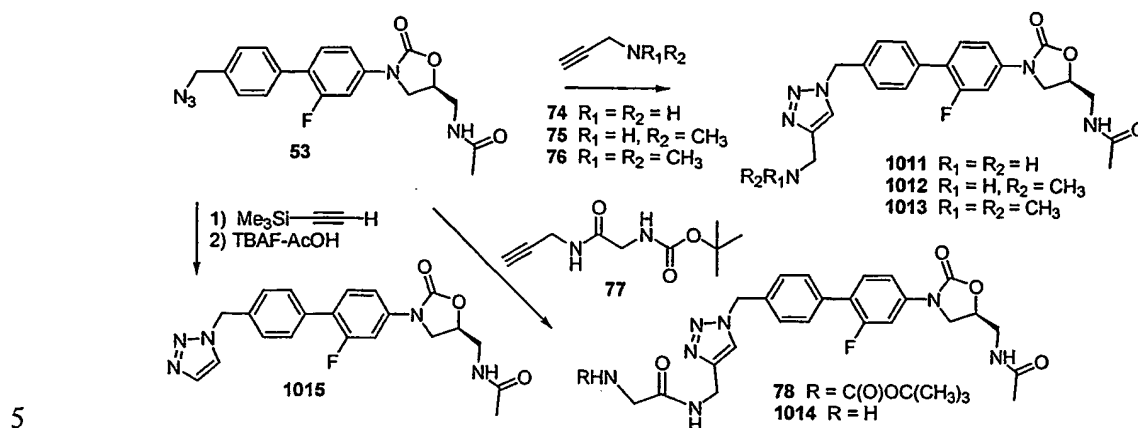
Compound **1010** was synthesized from mesylate **67** and imidazole in the same manner as described above for the synthesis of imidazole derivative **1008**. LCMS (ESI) m/z 427 ($\text{M} + \text{H}$)⁺.

Example 5 - Synthesis of Compounds **1011-1015**

Scheme 5 illustrates the synthesis of compounds **1011-1015**. The cycloaddition of azide **53** with alkynes **74-76** afforded triazoles **1011-1013** respectively. The cycloaddition of

azide **53** with alkyne **77** gave BOC-protected intermediate **78** which was subsequently cleaved to provide derivative **1014**. The cycloaddition of azide **53** with trimethylsilylacetylene, followed by desilylation, yielded triazole **1015**.

Scheme 5



Synthesis of triazole **1011**

A solution of azide **53** (0.10 g, 0.26 mmol) in propargyl amine **74** (0.50 mL) was treated with copper iodide (0.05 g, 0.26 mmol) and was stirred at 23 °C for 0.5 h. The reaction mixture was diluted with CH_2Cl_2 and MeOH and purified by flash chromatography and preparative TLC to afford **1011** as a brown solid (0.027 g; 24%). LCMS (ESI) m/z 439 ($\text{M} + \text{H}$)⁺.

10

Synthesis of triazole **1012**

A solution of azide **53** (0.10 g, 0.26 mmol) in N-methylpropargyl amine **75** (0.50 mL) was treated with copper iodide (5.00 mg, 0.026 mmol) and stirred at 23 °C for 12 h. The solvent was removed *in vacuo*, and the crude product was purified by preparative TLC to afford **1012** as a brown solid (0.038 g; 32%). LCMS (ESI) m/z 453 ($\text{M} + \text{H}$)⁺.

15

Synthesis of triazole **1013**

A solution of azide **53** (0.10 g, 0.26 mmol) in N, N-dimethylpropargyl amine **76** (0.056 mL, 0.520 mmol) was treated with copper iodide (5.00 mg, 0.026 mmol) and stirred at 23 °C for 12 h. The solvent was removed *in vacuo*, and the crude product was purified by flash chromatography to afford **1013** as a yellow film (0.073 g; 60%). LCMS (ESI) m/z 467 ($\text{M} + \text{H}$)⁺.

20

Synthesis of alkyne 77

A solution of propargyl amine **74** (0.34 mL, 5.0 mmol) in methylene chloride (25 mL) was treated with BOC-glycine (0.96 g, 5.5 mmol) and EDCI (1.1 g, 5.5 mmol) and stirred at 23 °C for 0.5 h. The reaction mixture was diluted with CH₂Cl₂, washed with 1.0 M HCl (aqueous), washed with saturated aqueous sodium bicarbonate (NaHCO₃), dried over Na₂SO₄, and the solvent evaporated *in vacuo* to afford alkyne **77** (0.51 g; 48%).

Synthesis of triazole 1014

A solution of azide **53** (0.15 g, 0.39 mmol) in THF (2 mL) was treated with alkyne **77** (0.17 g, 0.78 mmol) and copper iodide (7.00 mg, 0.039 mmol) and stirred at 23 °C for 16 h. The solvent was removed *in vacuo*, and the crude product was purified by flash chromatography to afford **78** as a white powder (0.16 g; 68%). LCMS (ESI) *m/z* 618 (M + Na)⁺.

A solution of **78** (0.15 g, 0.25 mmol) was treated with HCl (1.3 mL of 4.0 M solution in dioxane) and was stirred at 23 °C for 2 h. The solvent was removed *in vacuo*, and the residue twice redissolved in methylene chloride and evaporated to afford **1014** as a white film (0.14 g, 100%). LCMS (ESI) *m/z* 496 (M + H)⁺.

Synthesis of triazole 1015

A solution of azide **53** (0.75 mg, 2.0 mmol) in DMF (10 mL) was treated with trimethylacetylene (2.3 mL, 20 mmol) and was stirred at 90 °C for 12 h. The reaction mixture was cooled to 23 °C and the solvent was removed *in vacuo* to afford the expected silyl-substituted triazole as a brown foam (0.24 mg; 25%). LCMS (ESI) *m/z* 482 (M + H)⁺.

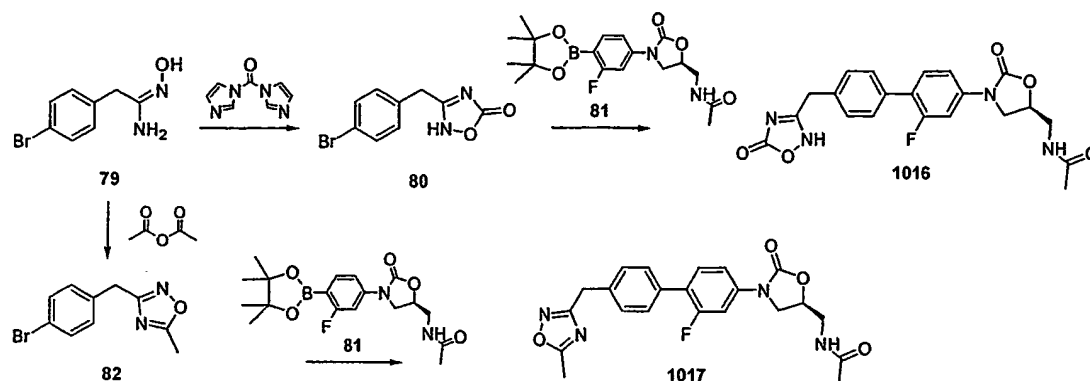
A solution of the above silyl-substituted triazole (0.050 g, 0.10 mmol) in THF (0.20 mL) was treated with acetic acid (6 µL, 0.10 mmol) and tetrabutylammonium fluoride (0.21 mL of 1.0 M solution in THF) and was stirred at 23 °C for 16 h. The reaction mixture was diluted with CH₂Cl₂, washed with water, dried (Na₂SO₄), and the solvent removed *in vacuo*. The crude product was purified to afford **1015** as a white powder (0.020 g; 47%). LCMS (ESI) *m/z* 432 (M + Na)⁺.

Example 6 - Synthesis of Compounds 1016-1017

Scheme 6 illustrates the synthesis of compounds **1016-1017**. Hydroxyamidine **79** was converted to bromide **80** which was subsequently coupled to boronate **81** to afford compound

1016. Hydroxyamidine 79 was transformed to oxadiazole 82, which was coupled to boronate 81 to afford compound 1017.

Scheme 6



5 Synthesis of hydroxyamidine 79

A solution of 4-bromophenylacetone nitrile (10 g, 54 mmol) in methanol (100 mL) was treated with sodium bicarbonate (2.2 g, 57 mmol) and hydroxylamine hydrochloride (4.0 g, 57 mmol) and refluxed for 1.5 h. Additional sodium bicarbonate (0.21 g, 5.4 mmol) and hydroxylamine hydrochloride (0.38 g, 5.4 mmol) were added, and the reaction mixture was refluxed for 12 h. The reaction mixture was cooled to 23 °C and the solvent removed *in vacuo* to afford hydroxyamidine 79 as a blue powder (4.0 g; 34%).

Synthesis of bromide 80

A solution of hydroxyamidine 79 (0.20 g, 0.91 mmol) in 1,4-dioxane (1 mL) was treated with 1,1'-carbonyldiimidazole (0.18 g, 1.1 mmol) and diazabicycloundecene (DBU, 0.15 mL, 0.97 mmol) and stirred at 105 °C for 1 h. The reaction mixture was diluted with water and extracted with ethyl acetate. The water layer was treated with 1.0 M HCl (aqueous) until the pH was 2, and then extracted with ethyl acetate. The organic layer was dried over Na₂SO₄, and the solvent removed *in vacuo* to afford bromide 80 as a yellow powder (0.11 g; 49%).

20 Synthesis of boronate 81

A suspension of *N*-[3-(3-fluoro-4-iodo-phenyl)-2-oxo-oxazolidin-5-ylmethyl]acetamide 62 (20.0 g, 52.8 mmol) in anhydrous 1,4-dioxane (130 mL) was treated with 4,4,5,5-tetramethyl-[1,3,2]dioxaborolane (10.2 g, 11.6 mL, 80.0 mmol) and triethylamine (16.0 g, 22.4 mL, 158.4 mmol) at room temperature, and the resulting reaction mixture was degassed three times under

a steady stream of argon before being treated with dichloro[1,1'-bis(diphenylphosphino)ferrocene] palladium (II) ($\text{Pd(dppf)}_2\text{Cl}_2$, 1.32 g, 1.6 mmol, 0.03 equiv) at room temperature. The reaction mixture was then degassed three times again under a steady stream of argon before being heated to reflux for 7 h. When TLC and LCMS showed that the reaction was complete, the reaction mixture was cooled down to room temperature before being treated with water (100 mL) and ethyl acetate (100 mL). The two layers were separated, and the aqueous layer was extracted with ethyl acetate (2 x 50 mL). The combined organic extracts were washed with water (2 x 50 mL) and saturated aqueous NaCl solution (50 mL), dried over magnesium sulfate (MgSO_4), and concentrated *in vacuo*. The residual brown oil was further dried *in vacuo* to afford the crude desired *N*-{3-[3-fluoro-4-(4,4,5,5-tetramethyl-[1,3,2]dioxaborolan-2-yl)-phenyl]-2-oxo-oxazolidin-5-ylmethyl} acetamide **81** (18.8 g, 20.0 g theoretical, 94%) as a brown solid which was of sufficient purity to be used in subsequent reactions.

Synthesis of compound 1016

A solution of boronate ester **81** (0.085 g, 0.220 mmol), bromide **80** (0.055 g, 0.220 mmol), and potassium carbonate (0.12 g, 0.90 mmol) in dioxane (1.4 mL), ethanol (0.46 mL) and water (0.46 mL) was degassed and treated with Pd(dppf)Cl_2 (6.0 mg, 6.7 μmol), degassed again, and heated at 80 °C for 1.5 h. The reaction mixture was diluted with CH_2Cl_2 and water, and the precipitate in the water layer was recovered by vacuum filtration to afford **1016** as a grey powder (0.034 g; 36%). LCMS (ESI) m/z 427 ($\text{M} + \text{H}$)⁺.

Synthesis of bromide 82

A solution of hydroxyamidine **79** (0.25 g, 1.1 mmol) in pyridine (5 mL) was cooled to 0 °C and treated with a solution of acetic anhydride (0.11 mL, 1.1 mmol) in pyridine (5 mL) and then stirred at 120 °C for 1.5 h. The reaction mixture was diluted with ethyl acetate, washed with 1.0 M HCl (aqueous), washed with saturated aqueous sodium bicarbonate, dried over Na_2SO_4 , and the solvent evaporated *in vacuo*. The crude product was purified by flash chromatography to afford bromide **82** as a clear film (0.10 g; 36%).

Synthesis of compound 1017

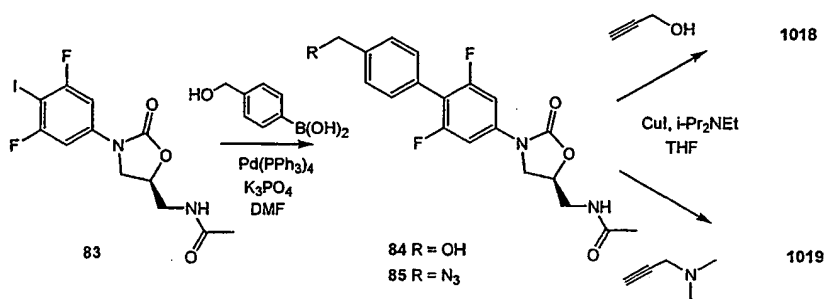
A solution of boronate ester **81** (0.15 g, 0.40 mmol), bromide **82** (0.10 g, 0.40 mmol), and potassium carbonate (0.22 g, 1.6 mmol) in dioxane (2.5 mL), ethanol (0.83 mL) and water (0.83 mL) was degassed and treated with Pd(dppf)Cl_2 (10.0 mg, 0.012 mmol), degassed again,

and stirred at 80 °C for 2 h. The reaction mixture was diluted with CH₂Cl₂ and washed with water. The water layer was extracted with 2 x CH₂Cl₂, dried over Na₂SO₄, and the solvent evaporated *in vacuo*. The crude product was purified by flash chromatography and preparative TLC to afford **1017** as a white powder (0.054 g; 32%). LCMS (ESI) *m/z* 425 (M + H)⁺.

5 Example 7 - Synthesis of Compounds 1018-1019

Scheme 7 illustrates the synthesis of compounds **1018-1019**. Known aryl iodide **83** was coupled to 4-hydroxymethylboronic acid to afford biaryl alcohol **84**. Alcohol **84** was converted to azide **85**, which was used in alkyne cycloaddition reactions to afford triazoles **1018** and **1019**.

10 Scheme 7



Synthesis of azide **85**

Known aryl iodide **83** (Gravestock, M.B., International Patent Application WO9910342) (1.00 g, 2.52 mmol) was dissolved in 6 mL DMF. 4-Hydroxymethyl-phenylboronic acid (0.461 g, 3.03 mmol) was added, followed by potassium phosphate (K₃PO₄, 0.804 g, 3.79 mmol) and Pd(PPh₃)₄ (0.292 g, 0.253 mmol). The mixture was degassed by evacuating the air from the flask, and refilling with argon (3 times), and then heated to 100°C for 4 hours. The mixture was allowed to cool and was then partitioned between ethyl acetate and water. The aqueous phase was extracted with ethyl acetate, and the combined organic phase was washed with brine, dried over MgSO₄, and evaporated. The residue was chromatographed on silica using a gradient mixture of methanol/methylene chloride (1% to 8%) to afford alcohol **84** (0.315 g, 0.838 mmol; 33%) as an ivory solid. An analytical sample was obtained by recrystallizing the material from methanol/methylene chloride/pentane. LCMS (ESI) *m/z* 377.

Alcohol **84** (0.889 g, 2.36 mmol) was suspended in 0.3 mL methylene chloride and 0.3 mL DMF. Triethylamine (0.66 mL, 4.74 mmol) was added, and the mixture was cooled to

0°C. Methanesulfonyl chloride (0.260 mL, 3.36 mmol) was added dropwise, and the mixture was stirred for 25 minutes. The mixture was then partitioned with ethyl acetate and water, and the organic layer was washed with brine, dried over MgSO₄, and evaporated. The residue was dissolved in 3 mL DMF, and sodium azide (0.384 g, 5.91 mmol) was added. The mixture was heated to 70°C for 4 hours. The reaction mixture was partitioned with ethyl acetate and water, and the organic layer was washed with brine, dried over MgSO₄, and evaporated. The residue was chromatographed on silica using a gradient mixture of methanol/methylene chloride (1% to 4%) to afford azide **85** (0.480 g, 1.20 mmol; 51%) as a tan solid. LCMS (ESI) *m/z* 402.

Synthesis of triazole 1018

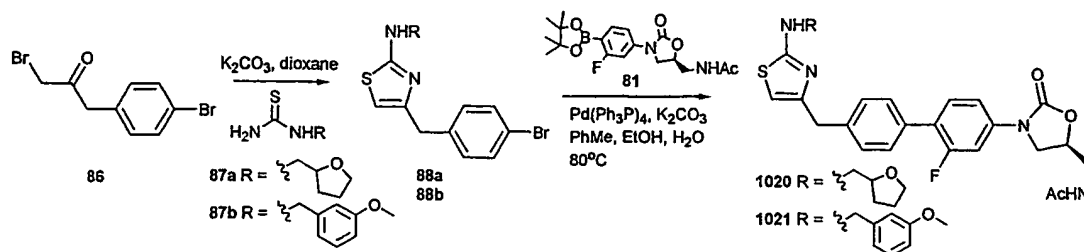
Azide **85** (0.084 g, 0.209 mmol) was dissolved in 0.7 mL THF and propargyl alcohol (25 µL, 0.400 mmol) was added, followed by Hunig's base (73 µL, 0.400 mmol) and copper(I) iodide (0.040 g, 0.210 mmol). The mixture was allowed to stir overnight at room temperature, and then was placed in a -20°C freezer for 2 days. The mixture was then partitioned with ethyl acetate and water, and the aqueous layer was extracted with ethyl acetate and then 2% methanol/methylene chloride. The combined organic layer was washed with brine, dried over MgSO₄ and evaporated. The residue was chromatographed on silica using a gradient mixture of methanol/methylene chloride (1% to 8%) to afford triazole **1018** (0.060 g, 0.131 mmol; 63%) as an ivory solid. LCMS (ESI) *m/z* 458.

Synthesis of triazole 1019

Azide **85** (0.135 g, 0.337 mmol) was dissolved in 1.5 mL THF and dimethyl-prop-2-ynyl-amine (72 µL, 0.674 mmol) was added, followed by *i*-Pr₂NEt (117 µL, 0.674 mmol) and copper(I)iodide (0.064 g, 0.337 mmol). The mixture was allowed to stir overnight at room temperature (the solvents evaporated overnight with positive pressure from argon gas). The residue was suspended in ethyl acetate and methylene chloride and filtered through celite. The pad of celite was washed with ethyl acetate and methylene chloride, and the combined organic washes were evaporated. The residue was chromatographed on silica using a gradient mixture of methanol/methylene chloride (0% to 14%) and the product obtained was triturated with methylene chloride and pentane. The tan solid was collected to afford triazole **1019** (0.072 g, 0.149 mmol; 44%). LCMS (ESI) *m/z* 485.

Example 8 - Synthesis of Compounds 1020-1021

Scheme 8 illustrates the synthesis of compounds **1020-1021**. Bromoketone **86** was subjected to alkylation with thioureas **87a** and **87b** to afford thiazoles **88a** and **88b** respectively. Coupling of **88a** and **88b** with boronate **81** yielded thiazoles **1020** and **1021**.

5 **Scheme 8****Synthesis of thiazole 88a**

Bromoketone **86** (0.29 g, 1.0 mmol) was dissolved in dioxane (10 mL). Thiourea **87a** (0.19 g, 1.2 mmol) and potassium carbonate (0.28 g, 2 mmol) were added sequentially and the resulting slurry stirred at $50^\circ C$ for 4 h. The mixture was cooled to room temperature, diluted with 100 mL CH_2Cl_2 , and washed with sat. aq. $NaHCO_3$, and brine. The aqueous washes were back-extracted with CH_2Cl_2 (2 x 50 mL). The combined organic extracts were dried over K_2CO_3 , filtered and concentrated *in vacuo* to afford **88a** as a yellow solid (0.32 g) which was used without further purification. LCMS (ESI) m/z 353 ($M + H$)⁺.

15 **Synthesis of thiazole 1020**

The crude aryl bromide **88a** obtained above (0.20 g, 0.56 mmol), boronate ester **81** (0.25 g, 0.66 mmol), and K_2CO_3 (0.14 g, 1.0 mmol) were combined with a 1:1:1 mixture of toluene, ethanol and water (2 mL each). The slurry was degassed by alternately applying high vacuum to the reaction mixture and flushing with dry argon. The reaction vessel was then sealed and heated in an $80^\circ C$ oil bath for 14 h. The reaction mixture was cooled to room temperature, diluted with 100 mL 9:1 CH_2Cl_2 /MeOH, and washed with water and brine (50 mL each). The aqueous washes were back-extracted once with 50 mL 9:1 CH_2Cl_2 /MeOH. The combined organic extracts were dried on K_2CO_3 , filtered, and concentrated *in vacuo* to afford 0.48 g of a brown solid which was purified by silica gel chromatography (25mm x 6" column eluted with 7:3 acetone/hexane) to yield **1020** as an off-white solid (0.17 g, 0.32 mmol). LCMS (ESI) m/z 525 ($M + H$)⁺.

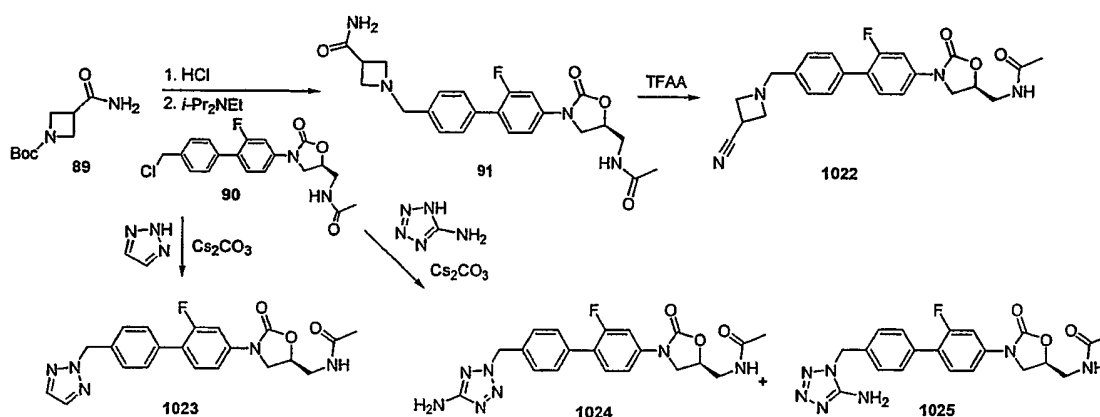
Synthesis of thiazole 1021

Compound **21** was synthesized according to the procedure described above for **1020**, using thiourea **88b** in place of **88a**. The reaction yielded **1021** as a white solid (0.12 g, 0.21 mmol). LCMS (ESI) m/z 561 ($M + H$)⁺.

5 Example 9 - Synthesis of Compounds 1022-1025

Scheme 9 illustrates the synthesis of compounds **1022-1025**. Azetidine **89** was deprotected and alkylated with chloride **90** to afford amide **91**. The amide of **91** was dehydrated with trifluoroacetic anhydride to produce nitrile **1022**. The alkylation of 1,2,3-triazole with benzyl chloride **90** gave triazole **1023**. Similarly, the alkylation of 5-aminotetrazole with benzyl chloride **90** yielded a mixture of tetrazole **1024** and tetrazole **1025**.

Scheme 9



Synthesis of chloride 90

N-[3-(2-fluoro-4'-hydroxymethyl-biphenyl-4-yl)-2-oxo-oxazolidin-5-ylmethyl]-acetamide **51** (3.0 g, 8.4 mmol) **51** was dissolved in CH₂Cl₂ (20 mL) and Hunig's base (2 mL). Methanesulfonyl chloride (1.4 mL, 12.6 mmol) was added dropwise and the resulting solution stirred at room temperature for 4 h. The mixture was poured into 100 mL sat. aqueous NaHCO₃ and extracted with CH₂Cl₂ (3 x 50 mL). The combined organic extracts were washed with brine, dried over MgSO₄, filtered, and concentrated to give 3.9 g of an oily yellow solid.

The crude material was purified by silica gel chromatography to give chloride **90** as an off-white solid (2.7 g, 7.2 mmol). LCMS (ESI) m/z 377 ($M + H$)⁺, 418 ($M + CH_3CN + H$)⁺, 440 ($M + CH_3CN + Na$)⁺.

Synthesis of amide 91

A solution of **89** (*J. Med. Chem.* 1993, 36, 801) (33 mg, 0.17 mmol) in CH₂Cl₂ (1.0 mL) was treated with 4.0 M HCl–dioxane (0.2 mL) and stirred at 23°C for 2 h. The reaction mixture was evaporated and the residue dissolved in DMF (1.0 mL) and treated with benzyl chloride **90** (63 mg, 0.17 mmol) and Hunig's base (0.17 mL, 1.0 mmol) and stirred at 60°C for 2 h. The reaction mixture was cooled to 23°C, diluted with H₂O (10 mL) and extracted with CH₂Cl₂ (4 × 25 mL), dried (Na₂SO₄) and evaporated. The crude residue was purified by preparative TLC (1% NH₄OH–10% MeOH–89% CH₂Cl₂) to afford **91** (36 mg; 50%) as a tan powder. LCMS (ESI) *m/z* 441.1 (M + H)⁺.

Synthesis of nitrile 1022

A solution of **91** (26 mg, 0.06 mmol) in CH₂Cl₂ (1.0 mL) was treated with pyridine (0.02 mL, 0.2 mmol) and trifluoroacetic anhydride (0.035 mL, 0.21 mmol) and stirred at 0°C for 1 h. The reaction mixture was directly purified by preparative TLC (1% NH₄OH–10% MeOH–89% CH₂Cl₂) to afford **1022** (6.0 mg; 24%) as a tan powder. LCMS (ESI) *m/z* 423.1 (M + H)⁺.

Synthesis of triazole 1023

A solution of **90** (0.19 g, 0.50 mmol) in DMF (2.0 mL) was treated with 1,2,3-triazole (0.058 mL, 1.0 mmol) and cesium carbonate (Cs₂CO₃, 0.33 g, 1.0 mmol) and stirred at 23°C for 16 h. The reaction mixture was diluted with H₂O (100 mL) and the resulting precipitate was isolated by filtration and purified by preparative TLC (10% MeOH–45% CH₂Cl₂–45% EtOAc) to afford **1023** (39 mg; 19%) as a white powder. LCMS (ESI) *m/z* 473.2 (M + CH₃CN + Na)⁺.

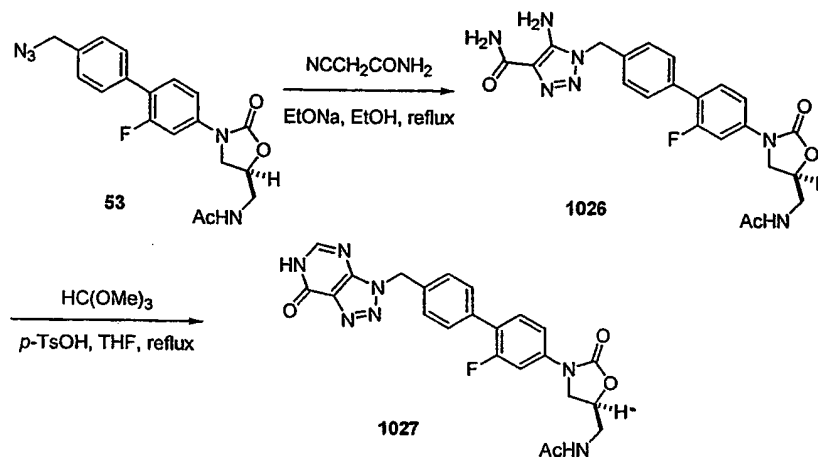
Synthesis of tetrazoles 1024 and 1025

A solution of **90** (0.19 g, 0.50 mmol) in DMF (2.0 mL) was treated with 5-aminotetrazole (87 mg, 1.0 mmol) and Cs₂CO₃ (0.33 g, 1.0 mmol) and stirred at 23°C for 12 h. The reaction mixture was diluted with H₂O (100 mL) and the resulting precipitate was isolated by filtration and suspended in 50 mL of a 1:1 mixture of CH₂Cl₂ and MeOH. The insoluble material (55 mg; 26%) was isolated by filtration and assigned the structure of **1024**. LCMS (ESI) *m/z* 426.1 (M + H)⁺. The soluble material was isolated by evaporation and purified by preparative TLC (1% NH₄OH–10% MeOH–89% CH₂Cl₂) to afford a white powder assigned the structure of **1025** (39 mg; 19%). LCMS (ESI) *m/z* 489.2 (M + CH₃CN + Na)⁺.

Example 10 - Synthesis of Compounds 1026 and 1027

Scheme 10 illustrates the synthesis of compounds **1026** and **1027**. Azide **53** was converted to triazole **1026**, which was then subsequently cyclized to compound **1027**.

Scheme 10



5

Synthesis of triazole 1026

A solution of azide **53** (383 mg, 1.0 mmol) in ethanol (4.0 mL) was treated with cyanoacetamide (101 mg, 1.2 mmol) and a solution of sodium ethoxide (21% wt solution in ethanol, 648 mg, 0.75 mL) at room temperature under N₂. The resulting reaction mixture was stirred for 10 min at room temperature before being warmed up to reflux for 2 h. When TLC showed that the reaction was complete, the reaction mixture was cooled down to room temperature before being treated with H₂O (10 mL). The white precipitate was then collected by filtration, washed with H₂O (2 x 10 mL), and dried *in vacuo* to afford the desired triazole **1026** (312 mg; 67%) as an off-white powder, which was of sufficient purity to be used directly in subsequent reactions. LCMS (ESI) *m/z* 468 (M + H)⁺.

15

Synthesis of compound 1027

A suspension of **1026** (165 mg, 0.353 mmol) in anhydrous THF (5 mL) was treated with *p*-toluenesulfonic acid monohydrate (34.2 mg, 0.18 mmol) and trimethyl orthoformate (374 mg, 0.386 mL, 3.53 mmol) at 25°C under N₂, and the resulting mixture was warmed up to reflux for 2 h. The solvents were removed *in vacuo*, and the residue was directly purified by column chromatography (5–10% MeOH/CH₂Cl₂ gradient elution) to afford the desired compound **1027** (42 mg; 25%) as a white powder. LCMS (ESI) *m/z* 478 (M + H)⁺.

20

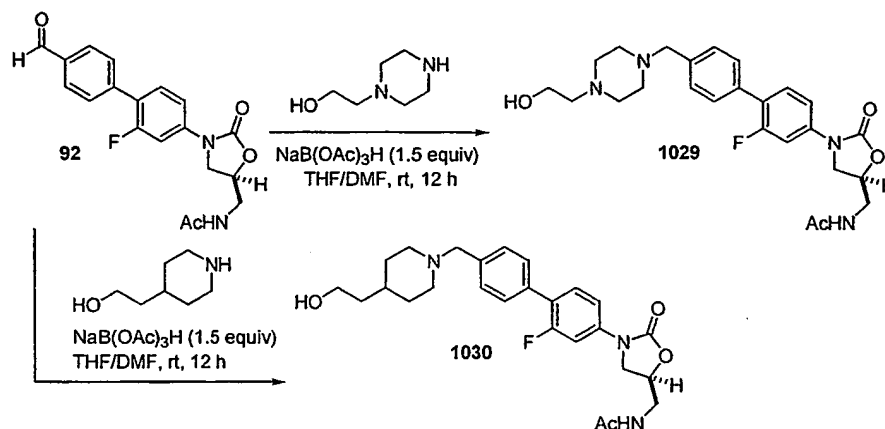
Example 11 - Synthesis of Triazole 1028

A suspension of azide **53** (124 mg, 0.324 mmol) in anhydrous 1,4-dioxane (5.0 mL) was treated with propargyl alcohol (182 mg, 0.19 mL, 3.24 mmol) at 25°C, and the resulting reaction mixture was warmed up to reflux for 12 h. When TLC and LCMS showed the reaction was complete, the reaction mixture was concentrated *in vacuo*, and the residue was directly purified by column chromatography (0–5% MeOH/CH₂Cl₂ gradient elution) to afford triazole **1028** (93.9 mg; 66%) as a pale-yellow solid. LCMS (ESI) *m/z* 440 (M + H)⁺.

Example 12 - Synthesis of Piperazine 1029 and Piperidine 1030

Scheme 11 illustrates the reductive amination chemistry used to synthesize **1029** and **1030**.

Scheme 11

**Synthesis of piperazine 1029**

A solution of aldehyde **92** (made from iodide **50** and 4-formylboronic acid in the same fashion as *N*-[3-(2-fluoro-4'-hydroxymethyl-biphenyl-4-yl)-2-oxo-oxazolidin-5-ylmethyl]-acetamide in Example 1) (180 mg, 0.5 mmol) and 2-piperidin-4-yl-ethanol (65 mg, 0.065 mL, 0.5 mmol) in anhydrous THF (4.0 mL) and anhydrous DMF (1.0 mL) was treated with sodium triacetoxyborohydride (160 mg, 0.75 mmol) at 25°C, and the resulting mixture was stirred at 25°C for 12 h. When TLC and LCMS showed the reductive amination reaction was complete, the reaction mixture was concentrated *in vacuo*. The residue was directly purified by flash column chromatography (0–5% MeOH-CH₂Cl₂ gradient elution) to afford piperazine **1029** (306 mg; 65%) as a colorless-oil, which solidified upon standing at room temperature *in vacuo*. LCMS (ESI) *m/z* 471 (M + H)⁺.

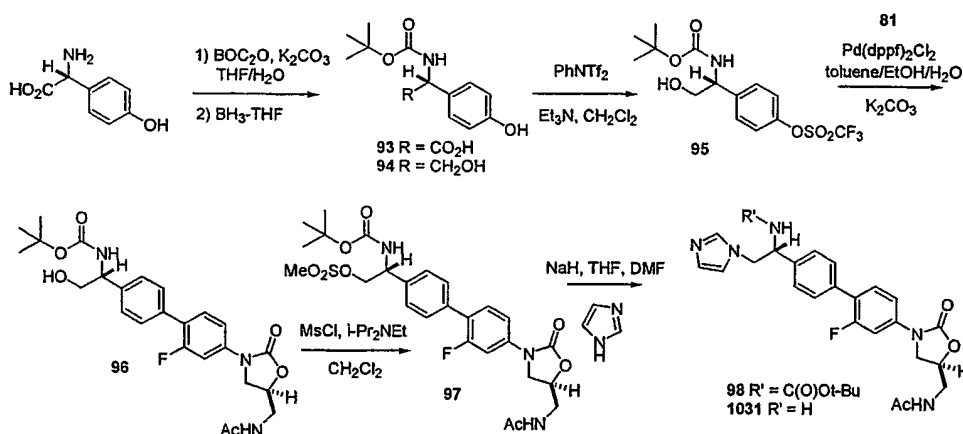
Synthesis of piperidine 1030

A solution of aldehyde **92** (356 mg, 1.0 mmol) and 2-piperazin-1-yl-ethanol (130 mg, 0.123 mL, 1.0 mmol) in anhydrous THF (8.0 mL) and anhydrous DMF (1.6 mL) was treated with sodium triacetoxyborohydride ($\text{NaB}(\text{OAc})_3\text{H}$, 318 mg, 1.5 mmol) at 25°C, and the resulting mixture was stirred at 25°C for 12 h. When TLC and LCMS showed the reductive amination reaction was complete, the reaction mixture was concentrated *in vacuo*. The residue was directly purified by flash column chromatography (0–5% $\text{MeOH}-\text{CH}_2\text{Cl}_2$ gradient elution) to afford piperidine **1030** (169 mg; 72%) as a colorless oil, which solidified upon standing at room temperature *in vacuo*. LCMS (ESI) m/z 470 ($\text{M} + \text{H}$)⁺.

Example 13 - Synthesis of Imidazole 1031

Scheme 12 depicts the synthesis of tetrazole derivative **1031**. D-*p*-Hydroxyphenyl-glycine was converted to triflate **95**, which was subsequently coupled to boronate **81** to afford alcohol **96**. Mesylation of **96**, followed by displacement with the anion of imidazole and deprotection of the BOC group yielded imidazole derivative **1031**.

Scheme 12



Synthesis of triflate 95

A solution of D-*p*-hydroxyphenylglycine (23.8 g, 142.3 mmol) and potassium carbonate (39.3 g, 284.6 mmol) in THF (200 mL) and H_2O (200 mL) was treated with di-*tert*-butyl dicarbonate (BOC_2O , 34.14 g, 156.6 mmol) at 25°C, and the resulting reaction mixture was stirred at 25°C for 2 h. When TLC and LCMS showed that the reaction was complete, the reaction mixture was treated with ethyl acetate (200 mL) and H_2O (200 mL). The two layers were separated, and the aqueous solution was extracted with ethyl acetate (200 mL), and the

combined organic extracts were discarded. The aqueous layer was then acidified with a 2 N HCl aqueous solution to pH 4 before being extracted with ethyl acetate (2 x 200 mL). The combined organic extracts were then washed with water (2 x 100 mL) and saturated aqueous NaCl solution (100 mL), dried over MgSO₄, and concentrated *in vacuo*. The residual white solids were further dried *in vacuo* to afford the crude desired acid **93** (36.5 g; 96%), which was of suitable purity for use in subsequent reactions.

A solution of acid **93** (4.005 g, 15 mmol) in anhydrous THF (20 mL) was treated dropwise with a 1 M solution of BH₃-THF in THF (30 mL, 30 mmol) at 0–5°C, and the resulting reaction mixture was stirred at 0–5°C for an additional 2 h. When TLC and LCMS showed that the reduction reaction was complete, the reaction mixture was treated with water (50 mL) and ethyl acetate (50 mL). The mixture was then stirred at 25°C for 30 min before being separated, and the aqueous layer was extracted with ethyl acetate (2 x 50 mL). The combined organic extracts were then washed with water (2 x 20 mL) and saturated aqueous NaCl solution (20 mL), dried over MgSO₄, and concentrated *in vacuo*. The residue was then directly purified by flash column chromatography (0-5% MeOH-CH₂Cl₂ gradient elution) to afford desired alcohol **94** (2.50 g; 66%) as a white powder which was of suitable purity for use in subsequent reactions.

A suspension alcohol **94** (670 mg, 2.65 mmol) in CH₂Cl₂ (10 mL) was treated with *N*-phenyltrifluoromethane sulfonamide (947 mg, 2.65 mmol) and triethylamine (535.3 mg, 0.74 mL, 5.3 mmol) at 25°C, and the resulting reaction mixture was stirred at 25°C for an additional 2 h. When TLC and LCMS showed that the reaction was complete, the reaction mixture was quenched with water (10 mL) and CH₂Cl₂ (20 mL). The two layers were then separated, and the aqueous layer was extracted with CH₂Cl₂ (2 x 20 mL). The combined organic extracts were then washed with water (2 x 10 mL) and saturated aqueous NaCl solution (10 mL), dried over MgSO₄, and concentrated *in vacuo*. The residue was then directly purified by flash column chromatography (0-5% MeOH-CH₂Cl₂ gradient elution) to afford triflate **95** (945 mg; 93%) as a white powder which was of suitable purity for use in subsequent reactions.

Synthesis of alcohol **96**

A solution of boronate **81** (2.162 g, 5.72 mmol) and triflate **95** (1.70 g, 4.4 mmol) in toluene (24 mL) was treated with solid potassium carbonate (1.82 g, 13.2 mmol), ethanol (8.0 mL) and H₂O (8.0 mL) at room temperature, and the resulting reaction mixture was degassed three times under a steady stream of argon before being treated with Pd(dppf)₂Cl₂ (184 mg,

0.22 mmol) at room temperature. The reaction mixture was then degassed three times again under a steady stream of argon before being warmed up to reflux for 2 h. When TLC and LCMS showed that the reaction was complete, the reaction mixture was cooled down to room temperature before being treated with water (20 mL) and ethyl acetate (20 mL). The two layers were separated, and the aqueous layer was extracted with ethyl acetate (2 x 20 mL). The combined organic extracts were washed with water (2 x 20 mL) and saturated aqueous NaCl solution (20 mL), dried over MgSO₄, and concentrated *in vacuo*. The residue was then purified by flash column chromatography (0-5% MeOH-CH₂Cl₂ gradient elution) to afford (1-{4'-[5-(acetylamino-methyl)-2-oxo-oxazolidin-3-yl]-2'-fluoro-biphenyl-4-yl}-2-hydroxyethyl)carbamic acid *tert*-butyl ester **96** (1.543 g; 72%) as yellow oil, which solidified upon standing at room temperature *in vacuo*.

Synthesis of mesylate **97**

A suspension of alcohol **96** (694 mg, 1.43 mmol) in anhydrous CH₂Cl₂ (10 mL) was treated with diisopropylethylamine (388 mg, 0.522 mL, 2.85 mmol) and methanesulfonyl chloride (196 mg, 0.132 mL, 1.71 mmol) at 0-5°C, and the resulting reaction mixture was stirred at 0-5°C for an additional 2 h. When TLC and LCMS showed that the reaction was complete, the reaction mixture was quenched with water (10 mL). The two layers were separated, and the aqueous layer was extracted with CH₂Cl₂ (2 x 10 mL). The combined organic extracts were washed with water (2 x 10 mL) and saturated aqueous NaCl solution (10 mL), dried over MgSO₄, and concentrated *in vacuo*. The residue was then purified by flash column chromatography (0-5% MeOH-CH₂Cl₂ gradient elution) to afford mesylate **97** (647 mg; 80%) as a pale-yellow solid, which was of suitable purity for use in subsequent reactions.

Synthesis of imidazole **98**

A solution of imidazole (41 mg, 0.6 mmol) in anhydrous THF (3 mL) was treated with NaH (60% oil dispersion, 29 mg, 0.72 mmol) at 0°C, and the resulting mixture was stirred at 0-5°C for 30 min before a solution of mesylate **97** (170 mg, 0.3 mmol) in anhydrous DMF (3.0 mL) was added. The resulting reaction mixture was then stirred at 0-5°C for 30 min before being gradually warmed up to room temperature for 12 h. When TLC and LCMS showed that the reaction was complete, the solvents were removed *in vacuo*, and the residue was directly purified by flash column chromatography (0-5% MeOH-CH₂Cl₂ gradient elution) to afford imidazole **98** (46 mg; 29%) as a yellow solid.

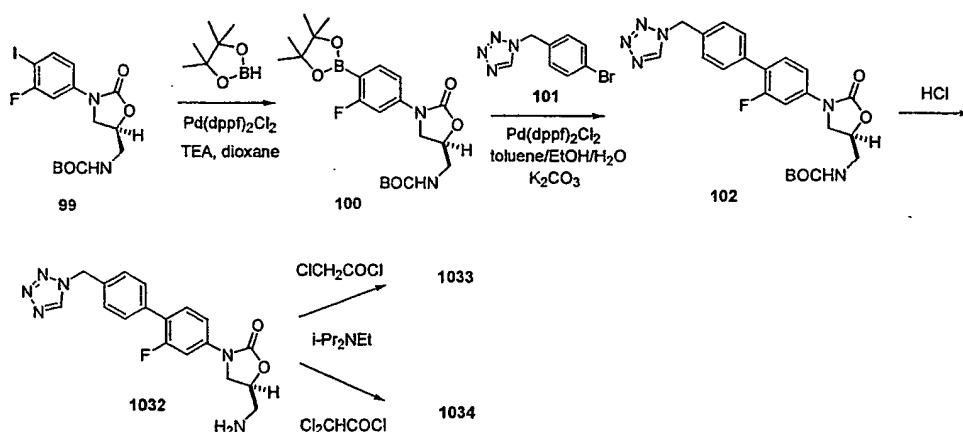
Synthesis of imidazole 1031

A solution of imidazole **98** (23 mg, 0.043 mmol) in MeOH (1.0 mL) was treated with a solution of 4 N HCl in 1,4-dioxane (3.0 mL), and the resulting reaction mixture was stirred at room temperature for 30 min. When TLC and LCMS showed that the reaction was complete, the solvents were removed *in vacuo*, and the desired *N*-{3-[4'-(1-amino-2-imidazol-1-yl-ethyl)-2-fluoro-biphenyl-4-yl]-2-oxo-oxazolidin-5-ylmethyl}acetamide hydrochloride **1031** (18.8 mg; 100%) was obtained as a yellow solid. LCMS (ESI) m/z 438 ($M + H$)⁺.

Example 14 - Synthesis of tetrazoles 1032-1034

Scheme 13 depicts the synthesis of tetrazole derivatives **1032-1034**. Iodide **99** was converted to boronate **100** which served as the coupling partner for bromide **101** to afford tetrazole **102**. Deprotection of **102** afforded tetrazole amine **1032**, which was subsequently acylated to afford tetrazole **1033** and **1034**.

Scheme 13



15 Synthesis of iodide 99

A solution of known 5-aminomethyl-3-(3-fluoro-4-iodo-phenyl)-oxazolidin-2-one (2.02 g, 6.0 mmol; *see* U.S. Patent Nos. 5,523,403 and 5,565,571) and potassium carbonate (1.66 g, 12.0 mmol) in THF (20 mL) and H_2O (20 mL) was treated with BOC_2O (1.334 g, 6.12 mmol) at 25°C, and the resulting reaction mixture was stirred at 25°C for 2 h. When TLC and LCMS showed the reaction was complete, the reaction mixture was treated with ethyl acetate (20 mL) and H_2O (20 mL). The two layers were separated, and the aqueous solution was extracted with ethyl acetate (20 mL), and the combined organic extracts were then washed with water (2 x 10 mL) and saturated aqueous NaCl solution (10 mL), dried over MgSO_4 , and concentrated *in*

vacuo. The residual white solids were further dried *in vacuo* to afford the crude, desired iodide 99 (2.40 g; 92%), which was of suitable purity for use in subsequent reactions.

Synthesis of boronate 100

A solution of iodide 99 (1.11 g, 2.55 mmol) in 1,4-dioxane (25 mL) was treated with
5 4,4,5,5-tetramethyl-[1,3,2]dioxaborolane (489 mg, 0.56 mL, 3.82 mmol) and triethylamine (772 mg, 1.07 mL, 7.65 mmol) at room temperature, and the resulting reaction mixture was degassed three times under a steady stream of argon before being treated with Pd(dppf)₂Cl₂ (107 mg, 0.13 mmol) at room temperature. The reaction mixture was then degassed three times again under a steady stream of argon before being warmed up to reflux for 6 h. When TLC and
10 LCMS showed that the reaction was complete, the reaction mixture was cooled down to room temperature before being treated with water (20 mL) and ethyl acetate (20 mL). The two layers were separated, and the aqueous layer was extracted with ethyl acetate (2 x 20 mL). The combined organic extracts were washed with water (2 x 20 mL) and saturated aqueous NaCl solution (20 mL), dried over MgSO₄, and concentrated *in vacuo*. The residual brown oil was
15 then purified by flash column chromatography (10-30% EtOAc-hexanes gradient elution) to afford boronate 100 (646 mg; 58%) as a brown oil, which solidified upon standing at room temperature *in vacuo* and was of suitable purity for use in subsequent reactions.

Synthesis of bromide 101

A solution of 4-bromobenzylamine hydrochloride (2.22 g, 10.0 mmol) in acetic acid (30
20 mL) was treated with triethyl orthoformate (2.964 g, 3.29 mL, 20.0 mmol) and sodium azide (2.30 g, 20.0 mmol) at room temperature, and the resulting reaction mixture was subsequently stirred at reflux for 12 h. When TLC and LCMS showed that the reaction was complete, the reaction mixture was cooled down to room temperature, and the cooled reaction mixture was poured into ice-water (100 mL). The precipitate was then collected by filtration, washed with
25 water (2 x 20 mL), and dried *in vacuo* to afford crude bromide 101 (460 mg; 19%) as a white solid, which was of suitable purity for use in subsequent reactions.

Synthesis of tetrazole 102

A solution of boronate 100 (658 mg, 1.5 mmol) and bromide 101 (300 mg, 1.25 mmol) in toluene (9.0 mL) was treated with solid potassium carbonate (621 mg, 4.5 mmol), ethanol
30 (3.0 mL) and H₂O (3.0 mL) at room temperature, and the resulting reaction mixture was degassed three times under a steady stream of argon before being treated with Pd(dppf)₂Cl₂

(52.3 mg, 0.063 mmol) at room temperature. The reaction mixture was then degassed three times again under a steady stream of argon before being warmed up to reflux for 3 h. When TLC and LCMS showed that the reaction was complete, the reaction mixture was cooled down to room temperature before being treated with water (10 mL) and ethyl acetate (20 mL). The two layers were separated, and the aqueous layer was extracted with ethyl acetate (2 x 10 mL). The combined organic extracts were washed with water (2 x 5 mL) and saturated aqueous NaCl solution (5 mL), dried over MgSO₄, and concentrated *in vacuo*. The residue was then purified by flash column chromatography (0-5% MeOH-CH₂Cl₂ gradient elution) to afford tetrazole 102 (357 mg; 61%) as a yellow oil, which solidified upon standing at room temperature *in vacuo*.

Synthesis of tetrazole 1032

A solution of tetrazole 102 (350 mg, 0.748 mmol) in EtOAc (5.0 mL) was treated with a solution of 4 N HCl in 1,4-dioxane (5.0 mL), and the resulting reaction mixture was stirred at room temperature for 30 min. When TLC and LCMS showed that the reaction was complete, the solvents were removed *in vacuo*, and the residue was treated with an aqueous sodium bicarbonate solution (10 mL) and EtOAc (15 mL). The mixture was stirred at room temperature for 30 min before the two layers were separated. The aqueous layer was extracted with EtOAc (10 mL), and the combined organic extracts were washed with H₂O (10 mL) and saturated aqueous NaCl solution (10 mL), dried over MgSO₄, and concentrated *in vacuo* to afford tetrazole amine 1032 (266 mg; 97%) as a pale-yellow solid. LCMS (ESI) *m/z* 369 (M + H)⁺.

Synthesis of tetrazole 1033

A suspension of tetrazole amine 1032 (74 mg, 0.2 mmol) in anhydrous CH₂Cl₂ (5.0 mL) was treated with diisopropylethylamine (52 mg, 0.07 mL, 0.4 mmol) and chloroacetyl chloride (34 mg, 0.024 mL, 0.3 mmol) at 0-5°C, and the resulting reaction mixture was stirred at 0-5°C for 2 h. When TLC and LCMS showed the reaction was complete, the reaction mixture was concentrated *in vacuo*. The residue was directly purified by flash column chromatography (0-5% MeOH-CH₂Cl₂ gradient elution) to afford tetrazole 1033 (43 mg; 48% yield) as a white solid. LCMS (ESI) *m/z* 445 (M + H)⁺.

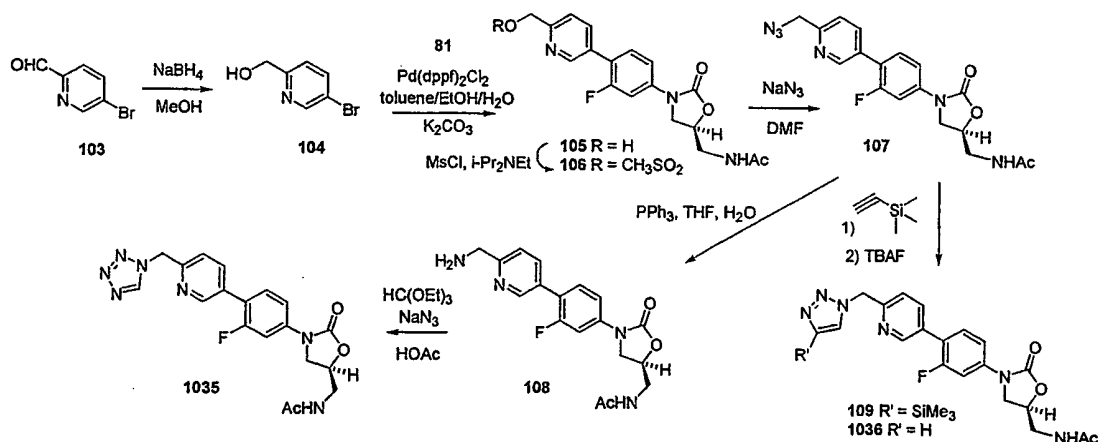
Synthesis of tetrazole 1034

A suspension of tetrazole amine **1032** (74 mg, 0.2 mmol) in anhydrous CH_2Cl_2 (5.0 mL) was treated with diisopropylethylamine (52 mg, 0.07 mL, 0.4 mmol) and dichloroacetyl chloride (44 mg, 0.029 mL, 0.3 mmol) at 0–5°C, and the resulting reaction mixture was stirred at 0–5°C for 2 h. When TLC and LCMS showed the reaction was complete, the reaction mixture was concentrated *in vacuo*. The residue was directly purified by flash column chromatography (0–5% MeOH- CH_2Cl_2 gradient elution) to afford tetrazole **1034** (41 mg; 43% yield) as a white solid. LCMS (ESI) m/z 479 ($\text{M} + \text{H}$)⁺.

Example 15 - Synthesis of compounds 1035 and 1036

10 Scheme 14 depicts the synthesis of tetrazole derivatives **1035** and **1036**. Aldehyde **103** was reduced to **104** which was coupled to boronate **81** to yield alcohol **105**. Mesylation of **105**, followed by displacement with sodium azide, yielded azide **107**. Reduction of **107** to amine **108** was followed by conversion to tetrazole **1035**. Cycloaddition of azide **107** with trimethylsilylacetylene, followed by desilylation, afforded triazole **1036**.

15 Scheme 14



Synthesis of aldehyde 103

A solution of 2,5-dibromopyridine (25 g, 105.5 mmol) in toluene (1.24 L) was cooled down to -78°C before being treated dropwise with a 2.5 M solution of *n*-BuLi in hexane (50.6 mL, 126.6 mmol) at -78°C under N_2 . The resulting reaction mixture was stirred at -78°C for 1 h before being treated with anhydrous DMF (11.6 g, 12.2 mL, 158.0 mmol) at -78°C . The reaction mixture was stirred at -78°C for an additional 1 h before being gradually warmed up to room temperature for 6 h. When TLC and LCMS showed that the reaction was complete, the